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(54) Title: NUCLEIC ACIDS, PROTEINS, AND ANTIBODIES

(57) Abstract: The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.







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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

## Nucleic Acids, Proteins, and Antibodies

- [1] This application refers to a "Sequence Listing" that is provided only on electronic media in computer readable form pursuant to Administrative Instructions Section 801(a)(i). The Sequence Listing forms a part of this description pursuant to Rule 5.2 and Administrative Instructions Sections 801 to 806, and is hereby incorporated in its entirety.
- The Sequence Listing is provided as an electronic file (PTZ15PCT\_seqList.txt, 1,891,228 bytes in size, created on January 13, 2001) on four identical compact discs (CD-R), labeled "COPY 1," "COPY 2," "COPY 3," and "CRF." The Sequence Listing complies with Annex C of the Administrative Instructions, and may be viewed, for example, on an IBM-PC machine running the MS-Windows operating system by using the V viewer software, version 2000 (see World Wide Web URL: http://www.fileviewer.com).

### Field of the Invention

[3] The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic

methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

#### Background of the Invention

- [4] The Human genome is estimated to contain roughly 100,000 genes, each of which plays an important function in sustaining life. Each of these roughly 100,000 genes encodes for a corresponding protein which can be classified based upon its structure and/or function. Some proteins are secreted, while other proteins reside either as membrane associated proteins or intracellularly. Although protein sequences vary substantially, many patterns and overall properties are shared, such as, for example, amino-terminal signal sequences.
- [5] Some proteins, for example secreted proteins, contain an amino-terminal signal sequence which facilitates protein transport. This amino-terminal signal sequence directs, or targets, the protein from its ribosomal assembly site to a particular cellular or extracellular location. Transport may involve any combination of several of the following steps: contact with a chaperone, unfolding, interaction with a receptor and/or a pore complex, addition of energy, and refolding. Moreover, an extracellular protein may be produced as an inactive precursor. Once the precursor has been exported, removal of the signal sequence by a signal peptidase activates the protein. Examples of some protein families that contain signal sequences include cytokines (chemokines) and hormones (growth and differentiation factors). Computer algorithms can be generated to identify amino-terminal signal sequences. Examples of computer programs designed to identify amino-terminal signal sequences include hidden Markov models (HMMs), statistical alternatives to FASTA and Smith Waterman algorithms, which have been used to find shared patterns, specifically consensus sequences (Pearson, W.R., and D.J. Lipman, PNAS, 85:2444-48 (1988); Smith, T.F., and M.S. Waterman, J. Mol. Biol., 147:195-97 (1981)). These algorithms are quite flexible in that they incorporate information from newly identified sequences to build even more successful patterns.
- Other families of proteins exist as membrane associated proteins. Examples of some of these membrane associated protein families include receptors (nuclear, 4

transmembrane, G protein coupled, and tyrosine kinase), protein kinases, phosphatases, neuropeptides and vasomediators, G proteins, ion channels (calcium, chloride, potassium, and sodium), proteases, transporter/pumps (amino acid, sugar, protein, metal and vitamin; calcium, phosphate, potassium, and sodium), matrix molecules (adhesion, cadherin, extracellular matrix molecules, integrin, and selectin), and regulatory proteins. Again, computer programs can aid in the discovery of these molecules. For example, Klein et al. have developed a method ("ALOM", also called as KKD) to detect potential transmembrane segments in polypeptides (Klein, P., et al., Biochim. Piophys. Acta., 815:468 (1985)). It attempts to identify the most probable transmembrane segment from the average hydrophobicity value over a range of amino acid residues. It predicts whether the segment is a transmembrane segment (INTEGRAL) or not (PERIPHERAL), and thus can suggest membrane association of a polypeptide.

- [7] Furthermore, some proteins function intracellularly, and can be identified by their structure and/or function. Computer algorithms can be adapted to aid in the identification of novel members of intracellular protein families. Examples of intracellular proteins include transcription factors, various classes of enzymes, Mitochondrial proteins, and signal transduction molecules.
- [8] Descriptions of some of these proteins (e.g., receptors, hormones, and matrix proteins) and diseases associated with their dysfunction follow.

#### Summary of the Invention

[9] The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

#### **Detailed Description**

#### **Tables**

[10] Table 1A summarizes some of the polynucleotides encompassed by the invention (including cDNA clones related to the sequences (Clone ID NO:Z), conting sequences (conting identifier (Contig ID:) and contig nucleotide sequence identifier (SEQ ID NO:X)) and further summarizes certain characteristics of these polynucleotides and the polypeptides encoded thereby. The first column provides the gene number in the application for each clone identifier. The second column provides a unique clone identifier, "Clone ID NO:Z", for a cDNA clone related to each contig sequence disclosed in Table 1A. The third column provides a unique contig identifier, "Contig ID:" for each of the contig sequences disclosed in Table 1A. The fourth column provides the sequence identifier, "SEQ ID NO:X", for each of the contig sequences disclosed in Table 1A. The fifth column, "ORF (From-To)", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence of SEQ ID NO:X that delineate the preferred open reading frame (ORF) that encodes the amino acid sequence shown in the sequence listing and referenced in Table 1A as SEQ ID NO:Y (column 6). Column 7 lists residues comprising predicted epitopes contained in the polypeptides encoded by each of the preferred ORFs (SEQ ID NO:Y). Identification of potential immunogenic regions was performed according to the method of Jameson and Wolf (CABIOS, 4; 181-186 (1988)); specifically, the Genetics Computer Group (GCG) implementation of this algorithm, embodied in the program PEPTIDESTRUCTURE (Wisconsin Package v10.0, Genetics Computer Group (GCG), Madison, Wisc.). This method returns a measure of the probability that a given residue is found on the surface of the protein. Regions where the antigenic index score is greater than 0.9 over at least 6 amino acids are indicated in Table 1A as "Predicted Epitopes". In particular embodiments, polypeptides of the invention comprise, or alternatively consist of, one, two, three, four, five or more of the predicted epitopes described in Table 1A. It will be appreciated that depending on the analytical criteria used to predict antigenic determinants, the exact address of the determinant may vary slightly. Column 8, "Tissue Distribution" shows the expression profile of tissue, cells, and/or cell line libraries which express the polynucleotides of the invention. The first number in column 8 (preceding the colon), represents the tissue/cell source identifier code corresponding to the key provided in Table 4. Expression of these polynucleotides was not observed in the other tissues and/or cell libraries tested. For those identifier codes in which

the first two letters are not "AR", the second number in column 8 (following the colon), represents the number of times a sequence corresponding to the reference polynucleotide sequence (e.g., SEQ ID NO:X) was identified in the tissue/cell source. Those tissue/cell source identifier codes in which the first two letters are "AR" designate information generated using DNA array technology. Utilizing this technology, cDNAs were amplified by PCR and then transferred, in duplicate, onto the array. Gene expression was assayed through hybridization of first strand cDNA probes to the DNA array. cDNA probes were generated from total RNA extracted from a variety of different tissues and cell lines. Probe synthesis was performed in the presence of <sup>33</sup>P dCTP, using oligo(dT) to prime reverse transcription. After hybridization, high stringency washing conditions were employed to remove nonspecific hybrids from the array. The remaining signal, emanating from each gene target, was measured using a Phosphorimager. Gene expression was reported as Phosphor Stimulating Luminescence (PSL) which reflects the level of phosphor signal generated from the probe hybridized to each of the gene targets represented on the array. A local background signal subtraction was performed before the total signal generated from each array was used to normalize gene expression between the different hybridizations. The value presented after "[array code]:" represents the mean of the duplicate values, following background subtraction and probe normalization. One of skill in the art could routinely use this information to identify normal and/or diseased tissue(s) which show a predominant expression pattern of the corresponding polynucleotide of the invention or to identify polynucleotides which show predominant and/or specific tissue and/or cell expression. Column 9 provides the chromosomal location of polynucleotides corresponding to SEQ ID NO:X. Chromosomal location was determined by finding exact matches to EST and cDNA sequences contained in the NCBI (National Center for Biotechnology Information) UniGene database. Given a presumptive chromosomal location, disease locus association was determined by comparison with the Morbid Map, derived from Online Mendelian Inheritance in Man (Online Mendelian Inheritance in Man, OMIM<sup>TM</sup>. McKusick-Nathans Institute for Genetic Medicine, Johns Hopkins University (Baltimore, MD) and National Center for Biotechnology Information, National Library of Medicine (Bethesda, MD) 2000. World Wide Web URL: http://www.ncbi.nlm.nih.gov/omim/). If the putative chromosomal location of the Query overlaps with the chromosomal location of a Morbid Map entry, an OMIM identification number is disclosed in column 10 labeled "OMIM Disease Reference(s)". A key to the OMIM reference identification numbers is provided in Table 5.

[11] Table 1B summarizes additional polynucleotides encompassed by the invention (including cDNA clones related to the sequences (Clone ID NO:Z), contig sequences (contig identifier (Contig ID:) contig nucleotide sequence identifiers (SEQ ID NO:X)), and genomic sequences (SEO ID NO:B). The first column provides a unique clone identifier, "Clone ID NO:Z", for a cDNA clone related to each contig sequence. The second column provides the sequence identifier, "SEQ ID NO:X", for each contig sequence. The third column provides a unique contig identifier, "Contig ID:" for each contig sequence. The fourth column, provides a BAC identifier "BAC ID NO:A" for the BAC clone referenced in the corresponding row of the table. The fifth column provides the nucleotide sequence identifier, "SEQ ID NO:B" for a fragment of the BAC clone identified in column four of the corresponding row of the table. The sixth column, "Exon From-To", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence of SEQ ID NO:B which delineate certain polynucleotides of the invention that are also exemplary members of polynucleotide sequences that encode polypeptides of the invention (e.g., polypeptides containing amino acid sequences encoded by the polynucleotide sequences delineated in column six, and fragments and variants thereof).

[12] Table 2 summarizes homology and features of some of the polypeptides of the invention. The first column provides a unique clone identifier, "Clone ID NO:Z", corresponding to a cDNA clone disclosed in Table 1A. The second column provides the unique contig identifier, "Contig ID:" corresponding to contigs in Table 1A and allowing for correlation with the information in Table 1A. The third column provides the sequence identifier, "SEQ ID NO:X", for the contig polynucleotide sequence. The fourth column provides the analysis method by which the homology/identity disclosed in the Table was determined. Comparisons were made between polypeptides encoded by the polynucleotides of the invention and either a non-redundant protein database (herein referred to as "NR"), or a database of protein families (herein referred to as "PFAM") as further described below. The fifth column provides a description of the PFAM/NR hit having a significant match to a polypeptide of the invention. Column six provides the accession number of the PFAM/NR hit disclosed in the fifth column. Column seven, "Score/Percent Identity", provides a quality score or the percent identity, of the hit disclosed in columns five and six. Columns 8 and 9, "NT From" and "NT To" respectively, delineate the polynucleotides in "SEQ ID NO:X" that encode a polypeptide having a significant match to the PFAM/NR database as disclosed in the fifth and sixth columns. In specific embodiments polypeptides of the invention comprise,

or alternatively consist of, an amino acid sequence encoded by a polynucleotide in SEQ ID NO:X as delineated in columns 8 and 9, or fragments or variants thereof.

[13] Table 3 provides polynucleotide sequences that may be disclaimed according to certain embodiments of the invention. The first column provides a unique clone identifier, "Clone ID", for a cDNA clone related to contig sequences disclosed in Table 1A. The second column provides the sequence identifier, "SEQ ID NO:X", for contig sequences disclosed in Table 1A. The third column provides the unique contig identifier, "Contig ID:", for contigs disclosed in Table 1A. The fourth column provides a unique integer 'a' where 'a' is any integer between 1 and the final nucleotide minus 15 of SEQ ID NO:X, and the fifth column provides a unique integer 'b' where 'b' is any integer between 15 and the final nucleotide of SEQ ID NO:X, where both a and b correspond to the positions of nucleotide residues shown in SEO ID NO:X, and where b is greater than or equal to a + 14. For each of the polynucleotides shown as SEQ ID NO:X, the uniquely defined integers can be substituted into the general formula of a-b, and used to describe polynucleotides which may be preferably excluded from the invention. In certain embodiments, preferably excluded from the invention are at least one, two, three, four, five, ten, or more of the polynucleotide sequence(s) having the accession number(s) disclosed in the sixth column of this Table (including for example, published sequence in connection with a particular BAC clone). In further embodiments, preferably excluded from the invention are the specific polynucleotide sequence(s) contained in the clones corresponding to at least one, two, three, four, five, ten, or more of the available material having the accession numbers identified in the sixth column of this Table (including for example, the actual sequence contained in an identified BAC clone).

Table 4 provides a key to the tissue/cell source identifier code disclosed in Table 1A, column 8. Column 1 provides the tissue/cell source identifier code disclosed in Table 1A, Column 8. Columns 2-5 provide a description of the tissue or cell source. Codes corresponding to diseased tissues are indicated in column 6 with the word "disease". The use of the word "disease" in column 6 is non-limiting. The tissue or cell source may be specific (e.g. a neoplasm), or may be disease-associated (e.g., a tissue sample from a normal portion of a diseased organ). Furthermore, tissues and/or cells lacking the "disease" designation may still be derived from sources directly or indirectly involved in a disease state or disorder, and therefore may have a further utility in that disease state or disorder. In numerous cases where the tissue/cell source is a library, column 7 identifies the vector used to generate the library.

Table 5 provides a key to the OMIM reference identification numbers disclosed in Table 1A, column 10. OMIM reference identification numbers (Column 1) were derived from Online Mendelian Inheritance in Man (Online Mendelian Inheritance in Man, OMIM. McKusick-Nathans Institute for Genetic Medicine, Johns Hopkins University (Baltimore, MD) and National Center for Biotechnology Information, National Library of Medicine, (Bethesda, MD) 2000. World Wide Web URL: http://www.ncbi.nlm.nih.gov/omim/). Column 2 provides diseases associated with the cytologic band disclosed in Table 1A, column 9, as determined using the Morbid Map database.

- [16] Table 6 summarizes ATCC Deposits, Deposit dates, and ATCC designation numbers of deposits made with the ATCC in connection with the present application.
- [17] Table 7 shows the cDNA libraries sequenced, and ATCC designation numbers and vector information relating to these cDNA libraries.
- [18] Table 8 provides a physical characterization of clones encompassed by the invention. The first column provides the unique clone identifier, "Clone ID NO:Z", for certain cDNA clones of the invention, as described in Table 1A. The second column provides the size of the cDNA insert contained in the corresponding cDNA clone.

#### **Definitions**

- [19] The following definitions are provided to facilitate understanding of certain terms used throughout this specification.
- [20] In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide. The term "isolated" does not refer to genomic or cDNA libraries, whole cell total or mRNA preparations, genomic DNA preparations (including those separated by electrophoresis and transferred onto blots), sheared whole cell genomic DNA preparations or other compositions where the art demonstrates no distinguishing features of the polynucleotide/sequences of the present invention.
- [21] As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence encoding SEQ ID NO:Y or a fragment or variant thereof; a nucleic acid sequence

contained in SEQ ID NO:X (as described in column 3 of Table 1A) or the complement thereof; a cDNA sequence contained in Clone ID NO:Z (as described in column 2 of Table 1A and contained within a library deposited with the ATCC); a nucleotide sequence encoding the polypeptide encoded by a nucleotide sequence in SEQ ID NO:B as defined in column 6 of Table 1B or a fragment or variant thereof; or a nucleotide coding sequence in SEQ ID NO:B as defined in column 6 of Table 1B or the complement thereof. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having an amino acid sequence encoded by a polynucleotide of the invention as broadly defined (obviously excluding poly-Phenylalanine or poly-Lysine peptide sequences which result from translation of a polyA tail of a sequence corresponding to a cDNA).

In the present invention, "SEQ ID NO:X" was often generated by overlapping [22] sequences contained in multiple clones (contig analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X is deposited at Human Genome Sciences, Inc. (HGS) in a catalogued and archived library. As shown, for example, in column 2 of Table 1A, each clone is identified by a cDNA Clone ID (identifier generally referred to herein as Clone ID NO:Z). Each Clone ID is unique to an individual clone and the Clone ID is all the information needed to retrieve a given clone from the HGS library. Furthermore, certain clones disclosed in this application have been deposited with the ATCC on October 5, 2000, having the ATCC designation numbers PTA 2574 and PTA 2575; and on January 5, 2001, having the depositor reference numbers TS-1, TS-2, AC-1, and AC-2. In addition to the individual cDNA clone deposits, most of the cDNA libraries from which the clones were derived were deposited at the American Type Culture Collection (hereinafter "ATCC"). Table 7 provides a list of the deposited cDNA libraries. One can use the Clone ID NO:Z to determine the library source by reference to Tables 6 and 7. Table 7 lists the deposited cDNA libraries by name and links each library to an ATCC Deposit. Library names contain four characters, for example, "HTWE." The name of a cDNA clone (Clone ID) isolated from that library begins with the same four characters, for example "HTWEP07". As mentioned below, Table 1A correlates the Clone ID names with SEQ ID NO:X. Thus, starting with an SEQ ID NO:X, one can use Tables 1, 6 and 7 to determine the corresponding Clone ID, which library it came from and which ATCC deposit the library is contained in. Furthermore,

it is possible to retrieve a given cDNA clone from the source library by techniques known in the art and described elsewhere herein. The ATCC is located at 10801 University Boulevard, Manassas, Virginia 20110-2209, USA. The ATCC deposits were made pursuant to the terms of the Budapest Treaty on the international recognition of the deposit of microorganisms for the purposes of patent procedure.

- In specific embodiments, the polynucleotides of the invention are at least 15, at least 30, at least 50, at least 100, at least 125, at least 500, or at least 1000 continuous nucleotides but are less than or equal to 300 kb, 200 kb, 100 kb, 50 kb, 15 kb, 10 kb, 7.5kb, 5 kb, 2.5 kb, 2.0 kb, or 1 kb, in length. In a further embodiment, polynucleotides of the invention comprise a portion of the coding sequences, as disclosed herein, but do not comprise all or a portion of any intron. In another embodiment, the polynucleotides comprising coding sequences do not contain coding sequences of a genomic flanking gene (i.e., 5' or 3' to the gene of interest in the genome). In other embodiments, the polynucleotides of the invention do not contain the coding sequence of more than 1000, 500, 250, 100, 50, 25, 20, 15, 10, 5, 4, 3, 2, or 1 genomic flanking gene(s).
- A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, or the complement thereof (e.g., the complement of any one, two, three, four, or more of the polynucleotide fragments described herein), the polynucleotide sequence delineated in columns 8 and 9 of Table 2 or the complement thereof, and/or cDNA sequences contained in Clone ID NO:Z (e.g., the complement of any one, two, three, four, or more of the polynucleotide fragments, or the cDNA clone within the pool of cDNA clones deposited with the ATCC, described herein), and/or the polynucleotide sequence delineated in column 6 of Table 1B or the complement thereof. "Stringent hybridization conditions" refers to an overnight incubation at 42 degree C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 μg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65 degree C.
- [25] Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions

include an overnight incubation at 37 degree C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH<sub>2</sub>PO<sub>4</sub>; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50 degree C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

- [26] Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.
- [27] Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone generated using oligo dT as a primer).
- [28] The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.
- [29] The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be

modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth. Enzymol. 182:626-646 (1990); Rattan et al., Ann. N.Y. Acad. Sci. 663:48-62 (1992)).

- "SEQ ID NO:X" refers to a polynucleotide sequence described, for example, in Tables 1Aor 2, while "SEQ ID NO:Y" refers to a polypeptide sequence described in column 6 of Table 1A. SEQ ID NO:X is identified by an integer specified in column 4 of Table 1A. The polypeptide sequence SEQ ID NO:Y is a translated open reading frame (ORF) encoded by polynucleotide SEQ ID NO:X. "Clone ID NO:Z" refers to a cDNA clone described in column 2 of Table 1A.
- [31] "A polypeptide having functional activity" refers to a polypeptide capable of displaying one or more known functional activities associated with a full-length (complete) protein. Such functional activities include, but are not limited to, biological activity,

antigenicity [ability to bind (or compete with a polypeptide for binding) to an antipolypeptide antibody], immunogenicity (ability to generate antibody which binds to a specific polypeptide of the invention), ability to form multimers with polypeptides of the invention, and ability to bind to a receptor or ligand for a polypeptide.

- [32] The polypeptides of the invention can be assayed for functional activity (e.g. biological activity) using or routinely modifying assays known in the art, as well as assays described herein. Specifically, one of skill in the art may routinely assay human polypeptides (including fragments and variants) of the invention for activity using assays as described in the examples section below.
- [33] "A polypeptide having biological activity" refers to a polypeptide exhibiting activity similar to, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention).
- [34] Table 1A summarizes some of the polynucleotides encompassed by the invention (including contig sequences (SEQ ID NO:X) and clones (Clone ID NO:Z) and further summarizes certain characteristics of these polynucleotides and the polypeptides encoded thereby.

Polynucleotides and Polypeptides of the Invention

TABLE 1A

Disease Reference(s): OMIM Cytologic Band AR061: 13, AR089: 10 S0050: 1 and S0260: 1. .0471: 1 and L0439: 1. Library code: count .0439: 1 and L0581: 1. Tissue Distribution AR089: 2, AR061: 1 (see Table IV for L0758: 2, S0010: 1, L0362: 3, L0794: 2, Library Codes) AR089: 1, AR061: L0769: 4, L0717: 3, H0622: 1, H0539: 1, H0624: 1, L0471: 1, AR061: 3, AR089: **Predicted Epitopes** Cys-118 to Thr-134 Pro-95 to Ser-103, Asp-17 to Phe-22, Pro-25 to Phe-36, Gly-51 to Gln-56, Arg-63 to Thr-68, Ser-75 to Phe-81. Tyr-84 to Trp-90, Ala-47 to Ser-62, Glu-70 to Pro-76. Gln-38 to Ser-44. Gln-9 to Ser-15. His-1 to Phe-10, Pro-1 to Arg-6, Pro-1 to Ile-13, NO: Y AA SEQ 335 336 562 338 561 337 (From-To) 247 - 516 77 - 232 2,-778 2 - 2501 - 243 2 - 346 ORF NO: X SEQ ID 238 237 14 12 13 Ξ 739539 906019 HAGDV32 1178626 Contig 1106393 699372 856958 Ä HFRBN59 Clone ID NO: Z HE2KJ64 HLICC37 Gene No: d n 4

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L0766: 3. L0774: 3.	L0775; 3, H0529; 2,	L0747: 2, L0756: 2,	L0777: 2, H0650: 1,	H0663: 1, S0442: 1,	S0358: 1, S0278: 1,	H0549: 1, H0318: 1,	H0052: 1, L0738: 1,	H0620: 1, H0014: 1,	H0355: 1, H0213: 1,	H0606: 1, S0448: 1,	S0142: 1, L0770: 1,	L0646: 1, L0773: 1,	L0651: 1, L0659: 1,	L0518: 1, L0663: 1,	H0547: 1, H0659: 1,	H0539: 1, L0748: 1,	L0750: 1, S0260: 1 and		AR089: 1, AR061: 1	H0617: 2, S0031: 2,	S0132: 1 and H0181: 1.		AR089: 1, AR061: 1	L0766: 4, H0038: 1,	H0616: 1, H0561: 1,	L0763: 1, H0521: 1,	L0750: 1, L0780: 1,
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L0758: 1 and L0595: 1.	AR061: 7, AR089: 5 H0255: 2 1 0493: 2	and L0662: 1.	·	•	AR089: 4, AR061: 2	L0539: 1 and H0553:	1.				AR089: 46, AR061: 9	H0617: 1						AR061: 3, AR089: 1	H0457: 5,-L0766: 5,	H0581: 2, H0090: 2,	H0521: 2, L0748: 2,	H0171: 1, H0656: 1,	S0212: 1, S0140: 1,	H0486: 1, H0156: 1,	L0471: 1, T0041: 1,
	Arg-14 to Arg-22, Pro-62 to Ala-79	Phe-106 to Arg-114,	Glu-120 to Gly-125.	,	Tyr-47 to His-53,	Lys-87 to Tyr-95,	Ser-110 to Ser-116,	Thr-124 to Ala-129,	Trp-146 to Arg-152.	Tyr-46 to His-52.	Pro-64 to Gly-71,	Lys-101 to Trp-106,	Glu-108 to Gly-116.	His-8 to Gly-18,	Pro-89 to Gly-96,	Lys-126 to Trp-131,	Glu-133 to Gly-141.	Val-30 to Leu-35,	Asn-65 to Leu-71,	Val-144 to Phe-149.	-	•	•		
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S0344: 1; S0426: 1,	L0387: 1, L0776: 1,	L0655: 1, L0367: 1,	L0792: 1, L0438: 1,	H0690: 1, H0539: 1,	H0436: 1, L0439: 1,	L0779: 1, L0780: 1,	L0755: 1 and H0422: 1.	AR089: 10, AR061: 4	L0759: 2 and H0593:		AR089: 8, AR061: 5	H0677: 54, L0604: 11,	S0366: 7, L0766: 6,	H0445: 6, H0543: 6,	H0556: 5, H0650: 5,	H0255: 5, L0770: 5,	L0655: 5, H0436: 5,	L0777: 5, L0485: 5,	H0657: 4, H0581: 4,	L0769: 4, L0761: 4,	L0747: 4, H0656: 3,	H0599: 3, H0196: 3,	H0373: 3, H0271: 3,	L0520: 3, L0546: 3,	H0423: 3, H0305: 2,	H0333: 2, L0623: 2,	H0457: 2, H0100: 2,
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18 - 478 567 Asp-27 to His-32, Gln-65 to Gly-76, Lys-80 to Ser-94, Pro-99 to Asn-104, Gly-126 to Lys-143, Pro-150 to Lys-143, Pro-150 to Lys-156, Glu-163 to Glu-175, Val-193 to Asp-204, Met-230 to Ser-263, Ala-278 to Gly-291, Pro-306 to Asn-320, Asn-328 to Lys-333, Glu-348 to Glu-355, Ile-358 to Asn-441, Ser-456 to Phe-463, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-535 to Glu-547. Thr-75 to Pro-94,								••																	
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18 - 478 567 Asp-27 to His-32, Gln-65 to Gly-76, Lys-80 to Ser-94, Pro-99 to Asn-104, Gly-126 to Lys-143, Pro-150 to Lys-143, Pro-150 to Lys-156, Glu-163 to Glu-175, Val-193 to Asp-204, Met-230 to Ser-263, Ala-278 to Gly-291, Pro-306 to Asn-320, Asn-328 to Lys-333, Glu-348 to Glu-355, Ile-358 to Asn-441, Ser-456 to Phe-463, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-484 to Lys-490, Glu-535 to Glu-547. Thr-75 to Pro-94,	l, H0522: 1, , L0750: 1, , L0731: 1, and L0584: 1																					٠			
18 - 478 567 Asp-27 to His-32, Gln-65 to Gly-76, Lys-80 to Ser-94, Pro-99 to Asn-104, Gly-126 to Lys-143, Pro-150 to Lys-143, Pro-150 to Lys-156, Glu-163 to Glu-175, Val-193 to Asp-204, Met-230 to Ser-263, Ala-278 to Gly-291, Pro-306 to Asn-320, Asn-328 to Lys-333, Glu-348 to Gly-355, Ile-358 to Asn-363, Glu-375 to Ser-381, Lys-390 to Arg-395, Lys-390 to Arg-395, Lys-490, Glu-484 to Lys-490, Glu-498 to Gly-507, Glu-535 to Glu-547.	H0518: J L0741: 1 L0752: 1 L0757: 1										•														
18 - 478 567		7 to His-32,	5 to Gly-76,	) to Ser-94,	to Asn-104,	26 to Lys-143,	0 to Lys-156,	53 to Glu-175,	3 to Asp-204,	30 to Ser-263,	78 to Gly-291,	6 to Asn-320,	28 to Lys-333,	18 to Glu-355,	3 to Asn-363,	75 to Ser-381,	30 to Arg-395,	33 to Asn-441,	6 to Phe-463,	34 to Lys-490,	98 to Gly-507,	35 to Glu-547.	to Ala-7,	5 to Pro-94,	Arg-111 to Gly-118,
18 - 478		Asp-2	Gln-65	Lys-80	Pro-99	Gly-12	Pro-15	Glu-16	Val-15	Met-23	Ala-27	Pro-30	Asn-32	Glu-34	Ile-358	Glu-37	Lys-35	Lys-43	Ser-45	Gln-48	Glu-49	Glu-52	Arg-1	Thr-75	Arg-1]
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Asp-122 to Gln-130.	His-1 to Glu-14,	Asp-26 to Lys-34,	Ser-47 to Lys-52,	Asn-97 to Gly-107,	Lys-123 to Gln-129,	Glu-215 to Asp-228,	Pro-245 to Glu-250,	Leu-255 to Glu-260,	Glu-275 to Gly-306.					•							٠	•					
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Glu-22 to Gly-35, Pro-37 to Thr-49.	Ser-40 to Ser-45, His-75 to Trp-81,	Ser-113 to Lys-128,	Pro-146 to Thr-154,	Asp-217 to Val-229,	Gly-261 to Gln-270,	Glu-313 to Thr-319,	Pro-346 to Leu-359,	Ala-378 to Ser-385,	Ser-388 to Asn-393,	Val-407 to Asp-418,	Ser-422 to Leu-428,	Thr-431 to Leu-441,	Leu-478 to Ala-489,	Gly-499 to Pro-522,	Glu-527 to Tyr-535,	Glu-540 to Arg-550,	Arg-560 to Arg-593,	Arg-625 to Ile-630,	Gin-642 to Tyr-649,	Lys-669 to Met-675,	Ala-687 to Thr-706,	Thr-734 to Asn-739.	Pro-78 to Lys-86,	Cys-88 to Leu-97,	Asp-100 to Ile-107,
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Pro-176 to Pro-181,	Arg-191 to Met-196,	Pro-200 to Arg-210,	Pro-246 to Ala-259,	Ser-271 to Glu-276,	Asp-298 to Trp-306,	Pro-332 to Ser-340.					•	-	•				-		•				-			. •	
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	Asn-7 to Asn-12.	Glu-19 to Asn-42, Ala-135 to Gly-140.	Leu-63 to Ser-76, Ala-141 to Pro-153, Pro-184 to Leu-194, Gln-235 to Gly-240, Pro-279 to Asp-288, Lys-296 to Pro-302.		Gly-9 to Gly-18, Arg-23 to Leu-28, Leu-65 to Ser-78, Ala-143 to Pro-155,
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Pro-186 to Leu-196, Gln-237 to Gly-242, Pro-281 to Asp-290, Lys-298 to Pro-304.	Glu-14 to Ala-21, Lys-51 to Ser-59, Ile-70 to Phe-75, Ala-107 to Arg-113, Thr-124 to Asn-131, Tyr-171 to Asn-176, Gln-187 to Asn-238, Ser-243 to Ile-248, Glu-265 to Ser-271, Pro-281 to Glu-298, Ser-309 to Met-316, Pro-321 to Pro-329, Gln-374 to Arg-381, Asp-390 to Cys-400.	
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Ser-86 to Ser-92.	Val-12 to Gln-17, Ala-75 to Arg-82, Lys-112 to Ile-117, Asn-179 to Trp-185, Asp-190 to Lys-209.	
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	Leu-10 to Pro-28.	Arg-7 to Tyr-12, Ser-46 to Lys-54, Gln-138 to Ile-147	Arg-13 to Ser-20, Asn-88 to Lys-94, Met-108 to Glu-113, Ala-154 to Phe-159, Arg-172 to Cys-202.
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	Leu-68 to Gln-77.	Gly-1 to Arg-7,	Ala-9 to Ser-15,	Ala-25 to Gly-30,	Gln-75 to Cys-84,	His-111 to Tyr-116.					-		-								-			Gly-10 to Lys-17.	Thr-10 to Ala-21,
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Gin-35 to Trp-45, Gly-54 to Leu-61.						Pro-30 to Gly-35.						Ser-6 to Glu-16,	Asp-33 to Lys-38,	Glu-71 to Phe-79,	Gln-120 to Glu-131,	Met-152 to Asp-159,	Ala-169 to Pro-174,	Leu-182 to Lys-201.		Pro-35 to Ser-43,	Glu-61 to Phe-69,	Gln-110 to Glu-120.	Gln-12 to Gln-17,	Arg-64 to Thr-69,	Ser-127 to Ser-132.
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	Thr-27 to Leu-32, Gly-42 to Gly-56, Ser-80 to Arg-90.	Phe-14 to Pro-20, His-23 to Ile-30, Ala-53 to Thr-58.	Thr-108 to Gly-115, Val-174 to Gly-181, Ala-205 to Gly-214, Pro-272 to Asn-282.	Pro-1 to Glu-13, Ser-22 to Lys-28, Gln-39 to Arg-50, Ser-111 to Asp-116.	
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	-		Pro-5 to Lys-22,	Arg-43 to Glu-51,	Arg-63 to Ala-71,	Asp-73 to Lys-79.		•								•				•								
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Gln-93 to Gln-100.	Thr-206 to Arg-212,	Gln-260 to Leu-269,	Arg-277 to Asp-284,	Arg-350 to Lys-357,	Arg-363 to Lys-378.	His-44 to Gly-49,	His-148 to Gly-154,	Lys-181 to Phe-204.		٠	•													:•		-	
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H0253: 1, H0309: 1, H0271: 1, H0039: 1, H0031: 1, H0087: 1, S0142: 1, L0763: 1, L0372: 1, L0644: 1, L0768: 1, L0375: 1, L0805: 1, L0653: 1, L0776: 1, L0653: 1, L0665: 1, H0672: 1, H0555: 1, L0740: 1, L0747: 1, L0779: 1, L0777: 1, L0731: 1, L0596: 1, S0276: 1 and H0552: 1.		AR089: 4, AR061: 2 H0543: 2	AR089: 7, AR061: 7 H0013: 2, H0560: 2, H0521: 2, H0624: 1, S6028: 1, S0038: 1, T0042: 1, L0475: 1, H0646: 1, S0426: 1,
Gln-310 to Leu-315.	Pro-3 to Gly-11, Gly-53 to His-63, Leu-70 to Lys-89, Met-99 to Thr-108.	Gly-1 to Lys-7.	Leu-39 to Arg-44.
	581	381	382
	3 - 428	1 - 195	169 - 726
	257	57	58
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L0766: 1, H0520: 1, H0519: 1, H0555: 1, H0542: 1 and S0424: 1.		AR089: 0, AR061: 0 H0497: 1 and T0039: 1.	AR089: 9, AR061: 6 H0635: 2, H0333: 1	and H0488: 1.		•			•	AR061: 4, AR089: 3	L0794: 4, S0360: 2,	H0553: 2, H0100: 2,	L0803: 2, L0741: 2,	L0745: 2, H0686: 1,	S0212: 1, S0418: 1,	S0420: 1, L0534: 1,	T0039: 1, H0013: 1,	H0575: 1, H0581: 1,	H0327: 1, H0428: 1,	T0006: 1, H0032: 1,	H0207: 1, S0002: 1,
	Leu-39 to Arg-44, Lys-178 to Asp-186.		Glu-20 to Ala-30, Glv-49 to His-62	Ser-75 to Gln-83,	Gly-148 to Gly-154,	Arg-158 to Ser-167,	Pro-169 to Pro-176,	Leu-213 to Val-222.	Gln-5 to Gly-10.	Arg-12 to Gln-23,	Asp-82 to Pro-88,	Gly-112 to Ala-120,	Arg-122 to Arg-127,	Gly-172 to Gly-186,	Val-212 to Gly-219,	Gly-242 to Gly-247,	Thr-253 to Ser-265.				
	582	383	384	•	•				583	385											
	288 - 691	1 - 402	681 - 1						1 - 252	2 - 826											
	258	59	09						259	61											
	911594	911588	1091714			•			894375	1154785											
		HSACD83	нндво53							HE8FD82								•			
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L0761: 1, L0499: 1, L0383: 1, L0519: 1, L0543: 1, L0789: 1, L0666: 1, H0144: 1, S0126: 1, H0658: 1, H0670: 1, H0436: 1, L0439: 1, L0786: 1, L0759: 1, L0596: 1 and L0592: 1.		AR089: 1, AR061: 0	H0046: 7, H0521: 7,	HUU52: 6, LU465: 6,	H0031: 5, H0624: 4,	S0358: 4, H0580: 4,	S0010: 4, S0346: 4,	H0551: 4, S0212: 3,	S0418: 3, S0007: 3,	H0437: 3, H0156: 3,	H0575; 3, H0457: 3,	L0471: 3, T0010: 3,	S0250: 3, H0328: 3,	H0644: 3, H0040: 3,	H0494: 3, S0344: 3,	S0002: 3, H0144: 3,	L0438: 3, H0520: 3,	S0152: 3, H0665: 3,	S0001: 2, H0402: 2,
	Ala-29 to Cys-34.	Ser-14 to Val-23,	Lys-76 to Ser-84,	Ser-102 to Leu-109,	Gln-119 to Cys-125,	Glu-177 to Thr-189,	Ala-221 to Phe-231.				-	• .				-	-		• •
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S0360: 2, S0046: 2,	H0393: 2, H0549: 2,	H0013: 2, H0069: 2,	H0318: 2, H0373: 2,	H0051: 2, S0214: 2,	H0553: 2, S0036: 2,	H0591: 2, H0038: 2,	H0616: 2, L0370: 2,	H0529: 2, H0519: 2,	H0539: 2, L0602: 2,	L0439: 2, L0591: 2,	S0026: 2, H0423: 2,	H0171: 1, L0615: 1,	S0040: 1, H0656: 1,	S0354: 1, H0329: 1,	H0369: 1, H0431: 1,	H0600: 1, H0586: 1,	H0559: 1, H0270: 1,	H0635: 1, H0427: 1,	H0590: 1, T0071: 1,	H0581: 1, N0006: 1,	H0123: 1, H0024: 1,	L0146: 1, H0014: 1,	S0003: 1, H0615: 1,	H0039: 1, H0622: 1,	H0030: 1, H0124: 1,	H0598: 1, H0090: 1,	L0060: 1, H0272: 1,
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S0015: 1, H0561: 1, S0440: 1, H0649: 1, H0646: 1, H0649: 1, S0210: 1, L0520: 1, L0378: 1, L0666: 1, S0374: 1, L0565: 1, H0547: 1, S0146: 1, H0547: 1, L0752: 1, S0260: 1, H0707: 1, L0779: 1, L0588: 1, L0596: 1, L0588: 1, L0593: 1, L0594: 1, H0653: 1 and H0422: 1.	AR089: 30, AR061: 6 H0013: 2 and L0485: 1.	AR089: 6, AR061: 4 S3014: 2	AR089: 6, AR061: 5 L0766: 6, L0748: 3, L0779: 3, S0360: 2, H0545: 2, H0494: 2,
	Glu-3 to Phe-8, Lys-43 to Glu-48, Gly-62 to Pro-71. Glu-3 to Phe-8, Lys-43 to Glu-48, Gly-62 to Pro-71.	Ala-94 to Cys-100, Ser-126 to Val-136, Val-161 to Asn-166. Ala-94 to Cys-100.	Pro-5 to Lys-12, Pro-18 to Arg-37, Asn-56 to Gly-63, Ser-75 to Arg-83,
	387	388	389
	1 - 513	1 - 540	606 - 118
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Gly-147 to Gly-156.		Ala-8 to Gly-13, Gln-58 to Cys-67,	His-94 to Tyr-99, Ser-107 to Ala-112.	Glu-29 to Leu-37, Ser-47 to Glu-53,	Glu-87 to Gln-92,		•	
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.0471: 3, T0010	S0250: 3, H0328:	H0644: 3, H0040:	H0494: 3, S0344:	30002: 3, H0144:	.0438: 3, H0520:	S0152: 3, H0665:	S0001: 2, H0402:	S0360: 2, S0046: 3	10393: 2, H0549:	H0013: 2, H0069	10318: 2, H0373: 2,	H0051: 2, S0214: 2,	40553: 2,	H0591: 2, H0038:	10616: 2,	10529: 2,	10539: 2,	943	S0026: 2, H0423:	H0171: 1,	S0040: 1, H0656:	S0354: 1, H0329:	H0369: 1, H0431:	H0600: 1	H0559: 1, H0270:	H0635: 1, H0427:	H0500.
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	H0581:	123:	146:	S0003: 1	H0039: 1	030:	598:	960:	S0015:	30440:	H0646:	210:	.0378:	S0374:	H0547: ]	10436: ]	779:	30260:	.0596: 1	,0485:	.0593: 1	H0653:	.061:	0266	592:	14.	013:	766:
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1 07	, ,	3	.0759: 2, S0242:	S0424: 2, H0624:	S0040: 1; S0420:	.0005: 1, S0356:	10357: 1, H0052:	H0009: 1, H0570:	, H0038:	H0413: 1, T0069:	, H0494:	0369: 1, L0794:	, L0803:	,L0651:	, H0520:	10435: 1, H0658:	.40666: 1, H0214:	S0028: 1, L0439: 1	and [	0, A	2, H(	9: I.	9, A	3, HC	, H0	, H01	H05	, H03
30.7		7:2	59: 2	24: 2,	10: 1;	05: 1	57: 1	99: 1	30051: 1,	13: 1	70041: 1	59: 1	.0649: 1	.0650: 1, ]	19990	35: 1	66: 1	28: 1,	.0755: 1	:68	060	and L0439: 1	.19	.992	51:2	27:2	52: 2,	59: 2
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Pro-188 to Lys-195.	Arg-1 to Glu-8, Ser-249 to Glu-254.
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			•	Leu-88 to Pro-94,	His-164 to Pro-174.	•					-						_				٠.						
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	Ser-10 to Gly-15, Pro-20 to Ser-27,	Glu-34 to Gly-41, Ala-45 to Trp-50, Pro-79 to Gly-88.		Ile-30 to Gly-36, Thr-67 to Thr-72.	Gly-15 to Arg-21,	Pro-30 to Ser-35,	Ser-44 to Asp-51,	Pro-109 to Phe-115,	Glu-131 to Ser-139,	Arg-166 to Ser-179,	Gly-205 to Gly-215,	Ser-234 to Arg-252,	Arg-279 to Glu-288,	Leu-355 to Cys-362,	Glu-371 to His-376,	Thr-393 to Asp-401,	Arg-506 to Asn-512,	Asp-571 to Lys-578,	Pro-580 to Pro-592,
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L0779: 1, H0543: 1 and H0506: 1.			AR089: 15, AR061: 2	S0126: 4, H0135: 3,	H0494: 3, H0547: 3,	S0045: 2, H0550: 2,	H0545: 2, H0242: 2,	H0266: 2, H0551: 2,	H0653: 2, S0040: 1,	S0282: 1, S0358: 1,	S0376: 1, S0046: 1,	H0393: 1, S6022: 1,	H0549: 1, H0156: 1,	H0618: 1, H0253: 1,	H0123: 1, H0050: 1,	H0024: 1, H0014: 1,	H0252: 1, H0124: 1,	H0040: 1, H0623: 1,	S0370: 1, S0210: 1,	L0648: 1, L0518: 1,	S0374: 1, H0435: 1,	S0328: 1, S0152: 1 and	_		
Lys-601 to Leu-608.	Ser-9 to Asp-16, Pro-74 to Phe-80.	Lys-85 to Gly-91.	Ser-26 to Val-32,	Ala-60 to Trp-66.	1												•			•			•	Pro-6 to Arg-14,	Gly-97 to Asp-109,
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·				AR089:	H026	L0592	H0144	H0013	10766	H0539: 2, 1	L0777: 2, I	L0759	S0424	S0040:	10005	H0357	H000	S0051	H0413:	T0041	L0369: 1	10649:	10650: 1	JT0666	H0435
Pro-118 to Cys-123, Cys-135 to Ser-140.	Pro-9 to Arg-17, Glv-100 to Asp-112.	Pro-121 to Cys-126,	Cys-138 to Ser-143.	Ser-75 to Lys-80,	Arg-167 to Lys-172.														•						
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H0666: 1, H0214: 1,	S0028: 1, L0439: 1,	L0755: 1 and L0593: 1.	AR089: 1, AR061: 1	L0748: 15, L0439: 6,	H0457: 4, H0009: 3,	H0620: 3, L0438: 3,	S0212: 2, H0559: 2,	H0673: 2, H0690: 2,	H0265: 1, H0341: 1,	H0305: 1, S0418: 1,	50360: 1, 50045: 1,	S0140: 1, S0278: 1,	H0549: 1, T0109: 1,	H0069: 1, H0590: 1,	H0618: 1, T0048: 1,	H0581: 1, H0052: 1,	H0050: 1, L0185: 1,	H0271: 1, H0213: 1,	H0617: 1, H0040: 1,	H0551: 1, H0264: 1,	L0769: 1, L0638: 1,	L0761: 1, L0644: 1,	L0771: 1, L0649: 1,	L0657: 1, L0809: 1,	L0666: 1, L0665: 1,	H0519: 1, H0689: 1,	H0435: 1, S0044: 1,
-		•	Ser-15 to Cys-21,	Leu-52 to Ser-58,	Gly-161 to Glu-167,	Arg-282 to Arg-289,	Ser-340 to Gln-345,	Arg-375 to Gln-381,	Gly-392 to Ala-399,	Pro-401 to Trp-406.	•				•			-		•							•
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	Gly-5 to Arg-12,	lle-52 to Thr-61,	Val-85 to Gly-92,	Tyr-114 to Thr-121,	Lys-133 to Pro-138,	Thr-186 to Arg-192.	Arg-1 to Ser-14,	Glu-46 to Glu-51.		-		Ala-1 to Gly-12.			•		•							
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1: 1, S	4: 1, S	7: 1, E		3: 1, H			0: 1, L		_	<u>_</u>		0: 1, E	0: 1, E	5: 1, H	2: 1, L	3: 1, S	5: 1, L	5: 1.		1: 1,	21: 3,	0: 1, S	7: 1, E	6: 1, E	3: 1 an	9: 16	50: 1 a
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	Gln-102 to Pro-108.							,																	
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	Lys-93 to Gln-98,	Asp-141 to Leu-148,	Asn-166 to Pro-172,	Glu-174 to Gln-179,	Ser-187 to Lys-192,	Gln-221 to Gln-229,	Pro-239 to Asp-246.	•												,							
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																	Val-51 to Arg-56,	Ala-127 to Asp-133,	Val-147 to Glu-153.	Gly-8 to Cys-13,	Gln-38 to Met-48,	Arg-76 to Gln-82,	Cys-87 to Asp-94.	Cys-22 to Cys-31,	Leu-35 to Pro-54,	Gln-59 to Glu-73,	Arg-131 to Met-140,	Asn-149 to Arg-156,
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Ser-191 to Thr-196,	Leu-299 to Leu-313,	Glu-328 to Pro-336,	Val-393 to Asp-399,	Asn-454 to Asn-466.								٠.															
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Asp-151 to Arg-157.	Lys-1 to Ser-18,	Asn-49 to Glu-62,	Gln-67 to Ser-76,	Glu-84 to Thr-90,	Thr-104 to Pro-112,	Ser-148 to Arg-156,	Gly-184 to Thr-191,	Pro-203 to Glu-210,	Thr-234 to Ser-240.											Lys-1 to Ser-18,	Asn-49 to Glu-62,	Gln-67 to Ser-76,	Glu-84 to Thr-90,	Thr-104 to Pro-112.	Leu-35 to Lys-41,	Leu-61 to Glu-68,	Ser-153 to Gln-158,
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•	Ser-9 to Cys-21,	Asn-137 to Leu-142,	Gly-231 to Thr-236,	Arg-284 to Phe-291,	Asn-305 to Asp-313,	Ala-375 to Asn-383,	Cys-404 to Arg-411,	Val-456 to Glu-469,	Glu-516 to Leu-521,	Lys-572 to Tyr-588.	Met-26 to Asn-37,	Glu-42 to Gln-51,	Thr-68 to Ser-95,	Ala-97 to Lys-113,	Asp-156 to Val-161,	Val-208 to Asp-215,	Pro-217 to Ala-228.							2	,		-
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	L0351: 1, H0494: 1,	H0561: 1, H0641: 1,	S0422: 1, L0763: 1,	L0769: 1, L0667: 1,	L0646: 1, L0800: 1,	L0643: 1, L0644: 1,	L0771: 1, L0662: 1,	L0768: 1, L0386: 1,	L0533: 1, L0806: 1,	L0653: 1, L0657: 1,	L0664: 1, H0691: 1,	H0518: 1, H0521: 1,	S0404: 1, H0436: 1,	L0743: 1, L0777: 1 and	L0600: 1.	AR089: 38, AR061: 7	L0731: 2 and S0010: 1.	AR089: 3, AR061: 0	S0001: 1, S0050: 1 and	H0181: 1.	AR061: 243, AR089:	175	S0050: 1, S0144: 1,	S0052: 1 and S0028: 1.	AR089: 11, AR061: 2	L0766: 14, L0761: 3,	L0792: 3, L0779: 3,	L0717: 2, H0135: 2,
				•							· ·	-						Glu-51 to Phe-60,	Gln-63 to Gly-73,	Thr-85 to Lys-91.	Asp-77 to Lys-82.				Pro-1 to Lys-13,	Pro-20 to Lys-39,	Ala-46 to Thr-71,	Pro-112 to Gln-122,
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H0264: 2, L0809: 2, L0790: 2, L0790: 2, L0791: 2, L0666: 2, L0591: 2, S0134: 1, H0650: 1, H0650: 1, H0580: 1, H0580: 1, H0580: 1, H0581: 1, H0590: 1, H0581: 1, H0694: 1, H0633: 1, S0364: 1, H0633: 1, S0362: 1, L0770: 1, L0769: 1, L0777: 1, S0194: 1 and H0543	•	9: 4, A 6: 10, L
H0264: 2, L0790: 2, L0666: 2, S0134: 1, H0657: 1, H0580: 1, H0050: 1, H0050: 1, S0364: 1, L0351: 1, L0770: 1, L0770: 1, L0770: 1, L0804: 1, L0783: 1,	, r	AR089: L0766:
rg-151, r-194, r-194, r-194, r-249, ra-285, e-303, rg-337, rg-337, sn-358, sp-395, sp-395, rg-445, rs-532, rg-562, rs-589, rg-589,	10, -36, -68, n-119, rg-148, -161, sp-192.	1-20, . 3-39,
Gly-129 to Arg-151, Gly-159 to Ile-164, Ala-188 to Tyr-194, Asn-208 to Pro-217, Gly-237 to Thr-249, Gly-267 to Ala-285, Ser-292 to Phe-303, Lys-305 to Ala-319, Asp-330 to Arg-337, Lou-347 to Asn-358, Val-388 to Ala-378, Yal-390 to Asp-395, Ser-417 to Arg-445, Phe-449 to Leu-476, Ala-510 to Lys-532, Ser-546 to Glu-562, Lys-570 to Ser-589, Val-609 to Glu-623.	Glu-5 to Lys-10, Pro-17 to Lys-36, Ala-43 to Thr-68, Pro-109 to Gln-119, Gly-126 to Arg-148, Gly-156 to Ile-161, Ala-185 to Asp-192.	Thr-15 to Glu-20, Val-29 to Arg-39,
Gly-1 Gly-1 Gly-2 Gly-2 Gly-2 Gly-2 Gly-2 Ser-2, Leu-3 Leu-3 Val-3 Ser-4 Phe-4 Phe-4 Phe-4 Phe-5 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-2 Cly-3 Cly-2 Cly-2 Cly-2 Cly-2 Cly-3 Cly-2 Cly-3	Glu-5 Pro-1 Ala-4 Pro-1 Gly-1 Gly-1 Ala-1	Thr-1 Val-2
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Pro-58 to Arg-66,	Lys-95 to Phe-105,	Val-109 to Ala-114.																•		-						·	
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•	Pro-8 to Gly-26, Cys-54 to Cys-66, Gly-73 to His-85.	Ser-31 to Gly-43, Ser-45 to Gly-57.	Ser-37 to Gly-49, Ser-51 to Gly-63, Val-93 to Cys-98.
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	Leu-12 to Ser-19, Glu-108 to Ser-119, Ala-121 to Thr-128, Lys-139 to Ala-149, Arg-153 to Ala-161.	Gln-43 to Ser-49, Ala-60 to Gly-67.	Gln-43 to Ser-49, Ala-60 to Gly-67, Arg-141 to Pro-146.	Trp-14 to Asp-27.	Asp-15 to Leu-21, Ser-59 to His-66, Ile-159 to Tyr-164.
	432	433	602	434	435
	98 - 556	1 - 840	1 - 582	147 - 362	1 - 585
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		Ile-11 to Arg-20, Pro-56 to Ala-69, Ser-81 to Glu-86,	Asp-108 to Trp-116, Pro-142 to Gly-158, Cys-207 to Ser-213.					. •	
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S0046: 1, H0619: 1,	S0222: 1, H0592: 1,	H0497: 1, H0574: 1,	H0632: 1, L0586: 1,	1, H0318: 1,	10046: 1, H0572: 1,	H0024: 1, S0051: 1,	F0010: 1, H0083: 1,	S6028: 1, H0266: 1,	H0271: 1, T0023: 1,	1, H0031: 1,	1, S0366: 1,		$\overline{}$	H0268: 1, H0412: 1,	H0059: 1, S0386: 1,	H0560: 1, S0150: 1,	l, S0344: 1,	1, S0426: 1,	H0529: 1, L0369: 1,	1, L0763: 1,	1, L0667: 1,	.0646: 1, L'0641: 1,	1, L0768: 1,		1, L0806: 1,	1, L0809: 1,	.0787: 1, L0792: 1,
S0046:	S0222: ]	H0497:	H0632:	L0021: 1	H0046:	H0024:	T0010:	Se028:	H0271:	L0483:	H0673:	H0135: 1	H0038:	H0268:	H0059:	H0560:	S0144: ]	H0538:	H0529:	L0640: 1	L0637: 1	L0646:	L0626:	L0387: 1	L0632:	L0655: 1	L0787:
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L0663: 1, H0691: 1,	H0660: 1, H0648: 1,	H0672: 1, S0328: 1,	S0378: 1, S0044: 1,	S0188: 1, H0134: 1,	S3012: 1, S0390: 1,	S0028: 1, L0749: 1,	L0786: 1, L0779: 1,	L0755: 1, L0757: 1,	L0758: 1, L0608: 1,	H0665: 1, H0542: 1 and	S0384: 1.	AR061: 1, AR089: 1	H0549: 7, L0665: 6,	L0751: 6, L0439: 5,	H0620: 3, L0803: 3,	L07.77: 3, L0601: 3,	H0483: 2, H0486: 2,	H0309: 2, L0774: 2,	L0657: 2, L0659: 2,	L0809: 2, L0666: 2,	L0438: 2, H0520: 2,	H0658: 2, L0602: 2,	H0555: 2, H0624: 1,	H0686: 1, H0295: 1,	H0656: 1, S0282: 1,	H0255: 1, S0354: 1,	H0580: 1, H0619: 1,
		,		•		-						Pro-11 to Ala-17,	Pro-19 to Gly-27,	Cys-60 to Gln-71,	Arg-73 to His-83,	Pro-85 to Asn-92.			•	•,	,						
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H0618: 1, H0581:	S0049: 1, H0052:	H0562: 1, H0012:	H0083: 1, H0687:	S0250: 1, H0428:	.0483: 1, H0135:	S0038: 1, H0494:	.0640: 1, L0638:	L0771:	L0805	L0629:	L0789:	1, H0519:	, H0682:	H0521:	H0522: 1, H0696:	J0740: 1, L0779: 1	H0667: 1 and H0543:	AR089: 19, AR061:	L0770: 2; S0126:-2,	.0748: 2, L0740: 2	H0542: 2, S0420:	S0360: 1, H0575:	H0551: 1, L0766: 1	J0745: 1, L0731: 1 and		6, AR089:	L0803: 6, L0748: 6,
618: 1,	)49: 1,	562: 1,	083: 1,	250: 1,	483: 1,	338: 1,	540: 1,	.0637: 1,	.0662: 1,	.0655: 1,	.0368: 1,	.0663: 1,	10593: 1,	670: 1,	522: 1,	740: 1,	667: 1	1089: 1	3770: 2	748: 2,	542: 2,	360: 1,	551: 1,	745: 1,	H0543: 1.	AR061:	0803: 6
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L0439: 6, L0777: 5, L0438: 3, S0330: 3, L0749: 3, S0282: 2, H0615: 2, L0770: 2	L0646: 2, L0754: 2, L0750: 2, L0756: 2,	L0779: 2, S0031: 2, L0592: 2, L0485: 2,	S0035: 1, H0574: 1,	10109: 1, H0013: 1, H0244: 1, H0590: 1,	S0346: 1, H0421: 1,	H0052: 1, T0110: 1,	S0050: 1, S0003: 1,	H0644: 1, H0383: 1,	7	H0598: 1, L0370: 1,	L0598: 1, L0521: 1,	L0662: 1, L0804: 1,	L0774: 1, L0775: 1,	L0653: 1, L0518: 1,	L0788: 1, H0144: 1,	H0660: 1, H0704: 1,	L0742: 1, L0740: 1,	L0759: 1, H0217: 1 and	H0543: 1.	AR089: 9, AR061: 5	H0090: 2, H0419: 1,
				,				•										-		Gln-24 to Pro-43,	Gly-68 to Lys-74.
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H0483: 1, H0459: 1, S0045: 1, H0455: 1, H0642: 1, H0485: 1, H0239: 1, H0617: 1, T0042: 1, H0649: 1, H0641: 1, H0547: 1, S0044: 1, S0037: 1, L0742: 1, L0439: 1, L0755: 1 and H0543: 1.	AR089: 52, AR061: 12 H0556: 1, H0346: 1, S0358: 1, H0090: 1, T0042: 1, H0560: 1, S0052: 1, H0519: 1 and S0152: 1.	AR061: 1, AR089: 1 H0581: 2, H0556: 1 and H0538: 1. AR061: 15, AR089: 4 H0617: 41, L0754: 38, L0779: 38, L0758: 32, H0618: 17, H0483: 11, S0358: 10, L0775: 10, L0777: 7, H0484: 6,
	Lys-34 to Ala-42, Lys-71 to Leu-76, Arg-188 to Trp-193, Val-215 to Asn-220, Ser-269 to Gln-274, Leu-333 to Lys-341, Thr-354 to Lys-361, Thr-401 to Ile-407, Lys-419 to Arg-427.	Ser-9 to Asn-15, Ser-64 to Gln-69. Cys-1 to Arg-13, Pro-15 to Gly-21, Gly-54 to Ser-59.
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L0774: 6, L0776: 6,	L0748: 6, L0740: 6,	L0752: 6, H0253: 5,	H0181: 5, T0114: 4,	L0750: 4, L0780: 4,	L0755: 4, H0606: 3,	H0087: 3, L0769: 3,	L0764: 3, L0771: 3,	L0806: 3, H0295: 2,	S0354: 2, H0549: 2,	H0298: 2, H0590: 2,	H0510: 2, H0553: 2,	H0038: 2, H0494: 2,	H0509: 2, L0783: 2,	L0809: 2, L0789: 2,	L0665: 2, S0330: 2,	H0696: 2, L0747: 2,	L0596: 2, H0653: 2,	H0661: 1, S0376: 1,	H0282:1, H0331:1,	H0574: 1, H0575: 1, .	H0251: 1, H0263: 1,	H0204: 1, H0596: 1,	T0110: 1, H0597: 1,	H0327: 1, L0719: 1,	H0544: 1, H0545: 1,	H0178: 1, H0620: 1,	H0375: 1, H0188: 1,
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H0615: 1, H0622: 1, H0033: 1, H0424: 1, H0644: 1, L0640: 1, L0763: 1, L0761: 1, L0642: 1, L0773: 1, L0662: 1, L0773: 1, L0662: 1, L0766: 1, L0669: 1, L0766: 1, L0669: 1, L0762: 1, L0666: 1, L0625: 1, L0666: 1, L0632: 1, L0666: 1, L0632: 1, L0666: 1, L0643: 1, L0741: 1, L0743: 1, L0731: 1, L0786: 1, L0731: 1, L0601: 1 and H0423: 1.	AR089: 1, AR061: 0 S0294: 2, H0559: 1 and L0747: 1.	AR089: 3, AR061: 0 S0040: 2, H0547: 2, L0393: 1, H0013: 1, H0427: 1, T0110: 1, T0078: 1, S0364: 1, H0124: 1, H0551: 1,
	Pro-30 to Ala-37, Ala-40 to Arg-49, Ala-152 to Leu-163. Pro-28 to Ala-35, Ala-38 to Arg-47.	Pro-12 to Ala-17, Asp-23 to Phe-28.
	446	447
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H0100: 1 H0494: 1	110100.1,110174.1,	H0509: 1, H0555: 1 and	L0439: 1.	AR061: 0, AR089: 0	H0255: 1, H0305: 1,	S0044: 1, S0037: 1,	S0028: 1 and S0031: 1.	AR061: 1, AR089: 1	H0494: 2, H0544: 1,	S0051: 1, L0754: 1 and	H0542: 1.		•	AR061: 3, AR089: 2	H0618: 3, L0439: 3,	H0124: 2, L0771: 2,	L0766: 2, S0126: 2,	H0445: 2, H0265: 1,	H0253: 1, H0318: 1,	H0421: 1, H0052: 1,	H0197: 1, H0015: 1,	S6028: 1, H0266: 1,	H0380: 1, H0529: 1,	L0803: 1, H0144: 1,	L0352: 1, S0328: 1,	H0539: 1, S0378: 1,	H0134: 1, L0749: 1,	L0777: 1, L0758: 1 and
				<u>                                     </u>				Ser-9 to Glu-14,	Arg-22 to Arg-27.			Arg-164 to Arg-169.		Lys-10 to Arg-25,	Glu-40 to Ala-46,	Arg-174 to Ala-181,	Ala-202 to Gln-208.											
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				714160				602854				847391	970432	1171692												····	·	
				HSDGJ23				HHSAD81						HCEEZ56								•						
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L0595: 1.			_		AR061: 6, AR089: 4	S0045: 2, S0046: 1,	H0645: 1, H0013: 1,	H0575: 1, H0286: 1,	H0521: 1 and H0136: 1.	-		AR061: 4, AR089: 4	L0439: 4, L0418: 1,	S0010: 1, L0455: 1,	S0028: 1 and L0741: 1.						•				AR061: 8, AR089: 7	H0081: 2, H0549: 1,
	Lys-10 to Arg-25,	Glu-40 to Ala-46,	Arg-174 to Ala-181,	Ala-202 to Gln-208.	Leu-26 to Tyr-32,	Pro-108 to Gln-123.				Leu-26 to Tyr-32,	Pro-108 to Gln-123.	Gly-14 to Glu-32,	Pro-60 to Ala-70,	Thr-145 to Gly-153,	Ser-164 to Leu-169.	-		-						Phe-4 to Gly-12.	Ser-38 to Asp-46,	Leu-55 to Leu-60,
	209				451					809		452												609	453	
	3 - 1346				3 - 2189					3-2189		3 - 509						-						473 - 138	601 - 1134	
	283				127				•	284		128			-									285	129	
	971572			•	1189455		,			952123		951351						,						956281	1202534	
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H0069: 1, H0046: 1,	H0428: 1, H0553: 1, H0087: 1, H0529: 1	.0532: 1, H0521: 1 and	H0423: 1.			!	AR061: 1, AR089: 1	L0748: 11, L0749: 6,	.0779: 4, L0438: 2,	H0547: 2, L0747: 2,	.0777: 2, L0596: 2,	H0650: 1, H0013: 1,	HO581: 1, H0046: 1,	10009: 1, H0266: 1,	H0622: 1, T0042: 1,	S0002: 1, H0695: 1,	H0529: 1, L0762: 1,	.0769: 1, L0771: 1,	.0766: 1, L0376: 1,	.0809: 1, L0666: 1,	.0665: 1, H0658: 1,	H0648: 1, S0044: 1,	H0555: 1, H0187: 1,	.0750: 1, L0752: 1,	L0758: 1, H0343: 1,	S0026: 1, S0192: 1,
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Lys-73 to Glu-79.			•	Ser-1 to Trp-6,	Ser-10 to Glu-22,	Pro-112 to Ser-117	Gln-15 to Asn-20,	Met-59 to Gln-66.		٠.			<b>4</b>					٠.								
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S0194: 1, H0542: 1, H0543: 1 and H0423: 1.		AR089: 10, AR061: 3	H0560: 2, S0342: 1,	H0586: 1, L0471: 1,	H0644: 1, H0617: 1,	H0040: 1, H0641: 1,	H0529: 1, H0519: 1,	S0037: 1 and L0757: 1.			AR089: 0	S3010: 2, S0028: 1 and	S0260: 1.				-			AR089: 1, AR061: 1	S0028: 3, H0617: 2,	S0045: 1, H0181: 1,	H0383: 1 and S0144: 1.	
	Gln-15 to Asn-20, Met-59 to Asn-64	Thr-1 to Asp-7,	Gly-37 to Asn-44,	Arg-175 to Tyr-180,	Lys-190 to Pro-198,	Gln-204 to Leu-209.			Thr-1 to Asp-7,	Gly-37 to Asn-44.	Arg-58 to Glu-63,	Val-80 to Gly-87,	Arg-114 to Lys-119,	Ala-132 to Gly-137,	Val-140 to Asp-145,	Ala-173 to Pro-178.	Ala-25 to Thr-31,	Glu-58 to Arg-63,	Gln-82 to Arg-87.	Arg-1 to Gly-8,	His-33 to Glu-44,	Ala-57 to Gly-62,	Tyr-71 to Arg-77,	Pro-85 to Asn-93,
	611	455							612	•	456						613			457				
	29 - 496	79 - 813			4			!	089 - 09		835 - 284						82 - 468		. •	808 - 2142	•			
	287	131					,		288		132						289			133	•			
	574258	1199942						_	882335		1193050						730740			1228145				
		HOUHW83									HSLCB60							•		HSLFG64				
		121									122				•					123				

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					-	AR061: 3, AR089: 2	L0750: 2, H0024: 1,	H0039: 1, H0622: 1,	H0040: 1 and S0434: 1.	AR061: 1, AR089: 0	L0748: 11, L0749: 6,	L0779: 4, L0438: 2,	H0547: 2, L0747: 2,	L0777: 2, L0596: 2,	H0650: 1, H0013: 1,	H0581: 1, H0046: 1,	H0009: 1, H0266: 1,	H0622: 1, T0042: 1,	S0002: 1, H0695: 1,	H0529: 1; L0762: 1,	L0769: 1, L0771: 1,	L0766: 1, L0376: 1,	L0809: 1, L0666: 1,	L0665: 1, H0658: 1,	H0648: 1, S0044: 1,	H0555: 1, H0187: 1,
Asp-116 to Ser-122.	Arg-1 to Gly-8,	Als 57 to Glu 69	Tyr 71 to Arg 77	Pro-85 to Ash-93	Asp-116 to His-121.	Asp-40 to Asn-49,	Cys-65 to Gly-71.	·		Ser-6 to Thr-11.	•					•			•	-	•					
	614					458				459																
	1196 - 3					3 - 422				677 - 2566		-				,				•						•
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L0750: 1, L0752: 1, L0758: 1, H0343: 1, S0026: 1, S0192: 1, S0194: 1, H0542: 1, H0543: 1 and H0423: 1.	AR089: 40, AR061: 37 S0015: 1 and H0665: 1	,	AR089: 56, AR061: 55 S0001: 1, S0051: 1 and S0028: 1.	•	AR061: 2, AR089: 1 S0278: 1, H0031: 1, H0617: 1 and S0390: 1.	AR089: 1, AR061: 0 H0013: 1, S0028: 1 and S0260: 1.		AR089: 5, AR061: 2 S0356: 1, S0354: 1, S0358: 1, S0376: 1,	H0620: 1, H0023: 1,
	Phe-5 to Val-11, Ser-28 to Lys-35, His-119 to Gln-127.	Phe-5 to Val-11, Ser-28 to Lys-35, His-119 to Gln-127.			Ser-19 to Asp-32, Tyr-58 to Gly-67.	Val-33 to Tyr-44.	Arg-52 to Lys-57, Glu-67 to Ile-74.	Leu-29 to Pro-47, Pro-55 to Arg-60, Pro-99 to Gly-106,	Met-170 to Thr-177,
,	460	615	461	616	462	463	617	464	
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	136	291	137	292	138	139	293	140	
	1121865	960388	1105323	791608	963457	882817	883338	949568	
	126 HKMAC08		HSLHS93	<u>*</u>	HBGOT10	HSDJW73		HWMEQ37	
	126		127		128	129		130	

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H0039: 1 and H0593: 1.	AR089: 1, AR061: 1	S0050: 1, H0316: 1,	S0428: 1, H0694: 1 and	120021: 1.			AR061: 3, AR089: 2	H0542: 2, H0597: 1,	H0288: 1, H0124: 1,	H0264: 1, S0344: 1,	L0752: 1 and L0581: 1.				AR089: 18, AR061: 9	S0150: 1				AR089: 1, AR061: 0	H0316: 1	AR061: 3, AR089: 2	L0748: 12, L0749: 7,	L0766: 5, L0803: 4,	L0756: 4, L0769: 3,	L0666: 3, H0547: 3,
Glu-196 to Ser-207.	Arg-6 to Gly-14,	Cys-20 to Gly-27,	Leu-80 to Pro-86.		Pro-6 to Thr-15,	Asp-27 to Thr-35.	Arg-8 to Arg-14.					Pro-21 to Ser-27,	Arg-42 to Asp-49,	Arg-82 to Ser-90.	Arg-12 to Tyr-23,	Ser-31 to Pro-37,	Thr-42 to Ala-56,	Ile-122 to Lys-128.	Gly-36 to Thr-41.			Pro-1 to Gly-6,	Ile-40 to Lys-46.			
	465				618	٠	466					619		•	467				620	468		469				
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	Gln-2 to Ser-11,	Pro-52 to Gly-61,	Thr-68 to Gly-103,	Lys-114 to Ala-120,	Pro-122 to Arg-127,	Gly-136 to Thr-147,	Asn-150 to Arg-167.	Gly-1 to Ser-11,	Ser-18 to Ala-25,	Ser-70 to Cys-77,	Asp-89 to His-104.	Ser-65 to Cys-72,	Asp-84 to His-99,	Arg-107 to Asn-112.	Ile-45 to Arg-52,	Phe-77 to Pro-85.					•		
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Ala-15 to Arg-31, Ala-55 to Gly-62, Glu-122 to Gly-128, His-150 to Asn-155, Val-187 to Arg-195.	Pro-1 to Gly-7, Gln-52 to Cys-61, His-88 to Tyr-93.	Ser-1 to Arg-6, Arg-37 to Thr-43.
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										Pro-1 to Pro-8,	Phe-12 to Thr-17,	Lys-60 to Gly-66,	Lys-71 to Gly-77,	Pro-88 to Gln-97.						Pro-8 to Trp-15,	Gly-42 to Gly-51,	Thr-58 to Arg-69.					Ast	<u>5</u>
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			Gly-1 to Ala-6,	Ser-19 to Ser-27,	Phe-31 to Leu-55,	Glu-72 to His-79,	Asn-120 to Gly-126,	Arg-158 to Arg-163.	Ser-9 to Ser-17,	Phe-21 to Leu-45.		· 							-			,	•			- •	
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Arg-31 to Val-38, Arg-80 to Pro-90, Asn-183 to Val-190, Val-318 to Leu-327, Leu-329 to Leu-354, Gln-357 to Glu-367, Leu-373 to Glu-380, Arg-391 to Gly-396, Ser-444 to Ser-457.	Arg-31 to Val-38, Arg-80 to Pro-90.				Met-1 to Gly-17, Pro-22 to Gly-30, Glv-72 to His-82.	Leu-89 to Lys-95.		Asn-31 to Leu-38, Cys-53 to Cys-64, Gly-139 to Cys-144.	
531	646	532	647	648	533			534	
64 - 1440	64 - 390	519 - 313	358 - 185	239 - 457	219 - 1433			1 - 432	
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	Gly-1 to Pro-11, Ser-39 to Thr-53.	Met-77 to Asn-92.	Gly-12 to Gly-20, Ser-86 to Glu-94	Pro-103 to Pro-110.									•			
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S002		AR061: 1, AR089:	2, H00	.0766: 2, L.0659: 2,	H0624: 1, H0341: 1,	, \$035	S0046: 1, H0370: 1	10427: 1, H0545:	.0769: 1, L.0764: 1	0794: 1, L0651:	, L078	.0438: 1, H0658: 1,	H0539: 1, L0439: 1	and H	AR089: 10, AR061:	12, L0	.0747: 6, L0755: 6,	, L074	.0439: 5, L0731: 5,	, L066	.0659: 4, L.0740: 4,	H0052: 3, H0090: 3	10040: 3, L0666: 3	H0547: 3, S0330: 3,	H0521: 3, L0779: 3,	.0608: 3, S0276: 3,	S0356: 2, S0360: 2,
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		Gln-9	o Ala-	o Ala-	o Ser-	to Ser						•			o Cys	o Gln	o Asp	to Se	to GI	to Gi	to Le	to Asp					
		Thr-3 to Gln-9,	Phe-36 to Ala-41,	His-52 to Ala-63,	Ala-81 to Ser-100,	Pro-122 to Ser-134.									Gly-58 to Cys-64,	Lys-74 to Gln-81,	Thr-90 to Asp-99,	Met-113 to Ser-118,	et-144	Gln-166 to Gly-173,	Thr-180 to Leu-187,	Ser-246 to Asp-256.					
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H0580: 2, H0156: 2, H0318: 2, H0050: 2	28: 2, H0032	S0366: 2, H0316: 2,	H0591: 2, H0059: 2,	.0763: 2, L0774: 2,	.0775: 2, L0809: 2,	565: 2, L0438	H0659: 2, H0555: 2,	.0744: 2, 1.0754: 2,	746: 2, H0171	H0265: 1, S0040: 1,	556: 1, H034]	58: 1, S0376	S0132: 1, H0619: 1,	H0587: 1, H0333: 1	185: 1, H0486	H0013: 1, S0010:	H0421: 1, H0251: 1	.0471: 1, H0014: 1	H0510: 1, H0375:	128: 1, H0553	H0617: 1, S0036:	H0412: 1, S0386:	H0641: 1, H0652:	S0422: 1, H0529:	.0773: 1, L0766:	.0776: 1, L.0527: 1
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	H0144: 1, H0682:	, HO	S0328: 1, H0539:	S0152: 1, H0522:	30146: 1, S0404: ]	, H04	L0750: 1, L0752:	100	and			4, A	2 and													51,7
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										Ser-12 to Arg-23,	Arg-56 to Asp-62.	Glu-26 to Pro-35,	Glu-56 to Ser-62,	Val	Ser-77 to Thr-82,	Val	o Gli	o Prc	Glu-26 to Pro-35,	Glu-56 to Ser-62,	Gln-67 to Val-73,	Ser-77 to Thr-82,	Ala-90 to Val-104	o Gl	o Prc	Ala-38 to Thr-45,
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Ser-70 to Asp-77,	Ser-85 to Asp-90,	Asp-139 to Gly-145,	Ile-207 to Asp-213,	Arg-229 to Met-234,	Gly-259 to Ser-264,	Ile-281 to Ser-288,	Asp-337 to Leu-343,	Gln-369 to Ile-376,	Gly-429 to Ser-440,	Gln-448 to Val-456,	Gln-461 to Thr-474.	•			•					•					-		
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Ala-38 to Thr-45,	Ser-/0 to Asp-//,	Ser-85 to Asp-90,	Asp-139 to Gly-145,	Arg-189 to Asp-196.	Pro-10 to Trp-21,	Ala-70 to Thr-79,	Leu-105 to Arg-111.	•						Asn-16 to Ser-23,	Lys-53 to Val-61,	Leu-77 to Asp-89,	Leu-116 to Ala-121,	Glu-152 to Lys-168,	Arg-178 to Lys-183,	Asp-196 to Glu-203,	Glu-220 to Ser-233.	Asn-16 to Ser-23,	Lys-53 to Asp-60.	Leu-27 to Pro-34,	Pro-40 to Lys-51,	Asn-85 to Phe-90,	Arg-102 to Leu-140,
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18 - 644					159 - 683								159 - 563	161 - 886								226 - 861		1 - 780			
328					219								329	220								330		221			
911607					1153892								766126	1152275								966135		974684		-	
					HHFGD38									HVA0G11										HUVDR03			
	•				209									210										211			

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.0439: 4, L0731: 4,	H0624: 3, S0222: 3,	H0457: 3, H0051: 3,	.0770: 3, L0769: 3,	.0790: 3, L.0666: 3,	547:	L0757: 3,	.0759: 3, H0050: 2,	H0056: 2, S0210: 2,	1,0774: 2,	L0665: 2,	L0748: 2,	.0751: 2, S0242: 2,	H0556: 1, H0657: 1	H0341: 1, H0484:	418:	300: 1	370: 1	438:	592:	250:	042:	9004:	1, H0421:	083:	l, H0355:	H0266: 1, H0271:	169:
4, L0	3,50	3, HC	3, LO	3, LO	3, HO	3, LO	3, H0	2, S0	2, 1.0	2, 1.0	2, L0	2, S0	1, HC	1, HC	H0125: 1, S0418:	S0354: 1, S0300: 1	1, HO	1, H0	H0600: 1, H0592:	70039: 1, H0250:	H0427: 1, H0042:	H0575: 1, H0004:	1, H0	H0012: 1, H0083:		1, H0	H0622: 1, H0169:
3439:	3624:	3457:	7770:	3790:	)664:	0750: 3, I	759:	3056:	.0662: 2, ]	.0519: 2, I	H0519: 2,	751:	3556:	3341:	3125:	354:	278:	3392:	3600:	039:	3427:	3575:	H0581: 1	3012:	H0408: ]	)266:	)622:
		王	<u> </u>	<u> </u>	걸	크	<u> </u>	王	<u> </u>	<u> </u>	Ħ	<u> </u>	H	Ħ	Ħ	SC	<u>80</u>	Ħ	Ħ	7	Ħ	丑	Ħ	Ħ	Ħ	H	Ħ
Gly-145 to Asp-191,	Glu-219 to His-227.												•														
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-145	1-219																										
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H0135: 1, H0264: 1, H0272: 1, H0488: 1, H0412: 1, H0623: 1,	H0059: 1, H0625: 1, H0641: 1, S0426: 1,	` `	<del>_</del>	,	L0663: 1, S0428: 1,	H0701: 1, S0148: 1,	L0438: 1, H0520: 1,	H0659: 1, H0648: 1,	H0672: 1, S0328: 1,	S0380: 1, H0627: 1,	H0631: 1, S0028: 1,	L0744: 1, L0754: 1,		L0752: 1, S0434: 1,	L0605: 1, L0485: 1,	H0136: 1, S0192: 1,	H0543: 1, H0422: 1 and	S0412: 1.	AR089: 13, AR061: 4	L0750: 2, H0370: 1,	 AR061: 1, AR089: 1	S0048: 1 and T0010: 1.	
				•						•											His-3 to Leu-15,	Tyr-28 to Ala-34,	Gly-52 to Glu-57,
				•							•								546		547		
																			58 - 288		1968 - 229		
									v.		-			•					222		223		
											•				-				689811		954681		
				•										•					HUDAE29		HIBCJ89		
			-			•		٠											212		213		

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									-	.•															
	-	AR089: 1, AR061: 1 L0439: 4, T0010: 2 and H0038: 1.	AR089: 14, AR061: 6	H0693: 44, L0604: 5,	S0366: 4, L0805: 4,	H0637: 3, L0766: 3,	H0672: 3, H0549: 2,	H0271: 2, L0777: 2,	H0661: 1, H0580: 1,	H0173: 1, H0620: 1,	H0428: 1, S0364: 1,	L0641: 1, L0644: 1,	L0655: 1, L0809: 1,	L0791: 1, L0666: 1,	L0663: 1, T0068: 1,	H0576: 1, L0780: 1,	L0731: 1, L0584: 1 and	H0543: 1.			AR089: 14, AR061: 8	L0766: 3, L0805: 3,	L0659: 3, L0744: 3,	L0794: 2, L0776: 2,	L0665: 2, H0648: 2,
Ser-123 to Gly-136.		Phe-15 to Glu-24.	Pro-26 to Ala-38,	Lys-85 to Gly-97,	Tyr-120 to Glu-131,	Asp-158 to Leu-168,	Asn-187 to Gly-197,	Ser-204 to Asp-209.	1								-		Lys-51 to Gly-63,	Tyr-86 to Glu-97.	Thr-1 to Arg-7,	Asn-34 to Gly-41,	Thr-67 to Asn-75.		
	655	548	549			•			•					•			٠		959	٠	550				
	367 - 1713	29 - 403	3 - 1784		•							•	•.		,				103 - 591		3 - 317				
	331	224	225																332		226				
	963279	504158	1195806		·										-		•		702070	-	1150878	•			
		HIBEG40	HWBEG33				,				•										HWHKD22				
		214	215																		216				

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H0305: 1, H0586: 1, H0599: 1, H0428: 1, H0551: 1, L0763: 1, L0637: 1, L0662: 1, L0768: 1, L0803: 1, L0804: 1, L0806: 1, L0655: 1, L0661: 1, L0787: 1, S0374: 1, H0520: 1, L0740: 1, L0770: 1, L0756: 1, L0777: 1, L0752: 1,		AR089: 1, AR061: 0 S0052: 1 and S0028: 1.	AR061: 1, AR089: 1 L0776: 20. L0777: 9.	L0439: 6, L0438: 4,	LU	H0024: 2, L0415: 1,	H0586: 1, H0596: 1,	H0050: 1, S0050: 1,	H0373: 1, S0051: 1,	S6028: 1, H0188: 1,	S0386: 1, S0448: 1,
	Asn-48 to Gly-55, Thr-81 to Asn-89.		Ser-40 to Tyr-45, Ala-61 to Pro-71.	Gly-92 to Asp-98,	Ala-145 to Asp-151, Pro-197 to Cys-205,	Leu-224 to Gly-235, Gly-241 to Ala-254	Ser-256 to Asn-262,	Asp-279 to Glu-290,	Ser-296 to Gly-303,	Lys-340 to Arg-345,	Ile-347 to Tyr-354.
		551	552								
	152 - 508	1 - 282	1 - 1083								
	333	227	228								
	963626	765497	944511	····		•			•		
		HSLF041	HE9SE46								
		217	218								

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L0369: 1,	L0775: 1,	H0144: 1,	S0330: 1,	L0750: 1,	,0779: 1, L0755: 1,	S0260: 1,	L0608: 1 and		AR089: 18, AR061: 18	H0618: 1, H0253: 1,	H0012: 1, H0620: 1,	H0181: 1 and H0617: 1.	1, AR061: 1			· · · · · ·								-		
S0306: 1, L0369:	1,00074: 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1,	L0805: 1, H0144:	T0068: 1, S0330:	L0745: 1, L0750:	L0779: 1,	L0731: 1,	L0596: 1,	H0665: 1.	AR089: 1	H0618:	H0012: 1,	H0181: 1	AR089:			•	,									
	•					-			Thr-1 to Leu-7.		•			Met-1 to Lys-11,	Asp-96 to Ile-104,	Asn-127 to Ser-140,	Gln-185 to Arg-190,	Lys-221 to Ser-231,	Ala-254 to Val-262,	His-295 to Asp-300,	Leu-304 to Ser-323,	Ser-327 to Gln-333,	Ala-345 to Ser-354,	Ala-370 to Ser-384,	Thr-396 to Gly-402,	Leu-413 to Pro-423,
				•			••		553		•		554	658												
•	·								216 - 836				17 - 2389	144 - 2336									,	,		
				:					229				230	334			:		•			•				
,					•		-		864276				1227138	1056330								-				
					•			•	HTLDW37				HWAFG54												•	
					•	•		٠	219	,			220													

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			103850, 114835, 116800, 140100, 192090, 192090,
	•		16q22.1
	AR089: 31, AR061: 25	AR061: 7, AR089: 4 S0116: 2, H0510: 2, H0144: 2, H0521: 2, L0748: 2, H0556: 1, T0049: 1, H0580: 1, H0393: 1, H0587: 1, H0051: 1, H0375: 1, H0622: 1, H0448: 1, H0646: 1, S0002: 1, L0752: 1 and L0731: 1.	AR061: 1, AR089: 1 H0457: 8, H0255: 6, L0743: 4, H0650: 2, S0354: 2, H0581: 2, L0747: 2, H0341: 1, S0376: 1, H0580: 1, H0069: 1, H0042: 1,
Gly-432 to Val-438, Ser-478 to Phe-485, Arg-487 to Lys-506, Ser-528 to Ser-547, Asn-557 to Ala-566, Asp-586 to Glu-597, Glu-644 to Pro-656, Leu-663 to Arg-671, Ser-700 to Asp-707.			Arg-50 to Gln-56, Gly-109 to Glu-119, Gln-131 to Asp-137, Gly-149 to Gly-159, Leu-184 to Glu-218, Val-239 to Ile-245.
	555	556	557
-	3 - 410	3 - 752	1 - 1302
	231	232	233
	810433	921175	932448
-	HKAFS73	HTXJD74	HSIGQ50
	221	222	223

192090, 192090, 245900, 245900, 276600, 600223		
H0036: 1, H0590: 1, H0251: 1, H0085: 1, H0123: 1, H0687: 1, H0213: 1, H0135: 1, H0040: 1, H0646: 1, S0002: 1, H0593: 1, H0555: 1, L0748: 1, L0731: 1, L0758: 1, L0731: 1, L0758: 1, H0506: 1,	AR089: 1, AR061: 1 H0657: 1, S0376: 1, H0123: 1, H0428: 1, L0646: 1, L0662: 1, L0803: 1, L0659: 1, L0790: 1, L0791: 1, H0660: 1 and L0759: 1.	AR089: 34, AR061: 19 L0664: 2, H0483: 1, S0376: 1, L0762: 1, L0638: 1, L0771: 1, L0657: 1, L0783: 1, L0665: 1, H0658: 1, H0670: 1 and L0779: 1.
	Gly-40 to Gly-46, Gln-60 to Arg-69, Lys-84 to Trp-91, Leu-112 to Arg-118.	Ser-15 to Tyr-24, Met-47 to Tyr-56, Gly-127 to Ser-133.
·	558	559
	3 - 647	3-677
	234	235
	932607	971537
	HWWDY45	HNSMB24
	224	225

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AR089: 3, AR061: 2 6	L0774: 3, L0771: 2,	L0766: 2, L0779: 2,	S0376: 1, L0646: 1,	L0764: 1, L0666: 1,	L0748: 1, L0731: 1,	L0593: 1 and H0423: 1.
			•	•		
260					•	
691 - 662			-	_		
236					•	
946862	•	,	,			
226   HWLOU63   946862						
226						

The first column in Table 1A provides the gene number in the application corresponding to the clone identifier. The second column in Table 1A provides a unique "Clone ID NO:Z" for a cDNA clone related to each contig sequence disclosed in Table 1A. This clone ID references the cDNA clone which contains at least the 5' most sequence of the assembled contig and at least a portion of SEQ ID NO:X was determined by directly sequencing the referenced clone. The reference clone may have more sequence than described in the sequence listing or the clone may have less. In the vast majority of cases, however, the clone is believed to encode a full-length polypeptide. In the case where a clone is not full-length, a full-length cDNA can be obtained by methods described elsewhere herein.

- [36] The third column in Table 1A provides a unique "Contig ID" identification for each contig sequence. The fourth column provides the "SEQ ID NO:" identifier for each of the contig polynucleotide sequences disclosed in Table 1A. The fifth column, "ORF (From-To)", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence "SEQ ID NO:X" that delineate the preferred open reading frame (ORF) shown in the sequence listing and referenced in Table 1A, column 6, as SEQ ID NO:Y. Where the nucleotide position number "To" is lower than the nucleotide position number "From", the preferred ORF is the reverse complement of the referenced polynucleotide sequence.
- [37] The sixth column in Table 1A provides the corresponding SEQ ID NO:Y for the polypeptide sequence encoded by the preferred ORF delineated in column 5. In one embodiment, the invention provides an amino acid sequence comprising, or alternatively consisting of, a polypeptide encoded by the portion of SEQ ID NO:X delineated by "ORF (From-To)". Also provided are polynucleotides encoding such amino acid sequences and the complementary strand thereto.
- [38] Column 7 in Table 1A lists residues comprising epitopes contained in the polypeptides encoded by the preferred ORF (SEQ ID NO:Y), as predicted using the algorithm of Jameson and Wolf, (1988) Comp. Appl. Biosci. 4:181-186. The Jameson-Wolf antigenic analysis was performed using the computer program PROTEAN (Version 3.11 for the Power MacIntosh, DNASTAR, Inc., 1228 South Park Street Madison, WI). In specific embodiments, polypeptides of the invention comprise, or alternatively consist of, at least one, two, three, four, five or more of the predicted epitopes as described in Table 1A. It will be appreciated that depending on the analytical criteria used to predict antigenic determinants, the exact address of the determinant may vary slightly.

[39] Column 8 in Table 1A provides an expression profile and library code: count for each of the contig sequences (SEQ ID NO:X) disclosed in Table 1A, which can routinely be combined with the information provided in Table 4 and used to determine the tissues, cells, and/or cell line libraries which predominantly express the polynucleotides of the invention. The first number in column 8 (preceding the colon), represents the tissue/cell source identifier code corresponding to the code and description provided in Table 4. For those identifier codes in which the first two letters are not "AR", the second number in column 8 (following the colon) represents the number of times a sequence corresponding to the reference polynucleotide sequence was identified in the tissue/cell source. Those tissue/cell source identifier codes in which the first two letters are "AR" designate information generated using DNA array technology. Utilizing this technology, cDNAs were amplified by PCR and then transferred, in duplicate, onto the array. Gene expression was assayed through hybridization of first strand cDNA probes to the DNA array. cDNA probes were generated from total RNA extracted from a variety of different tissues and cell lines. Probe synthesis was performed in the presence of <sup>33</sup>P dCTP, using oligo(dT) to prime reverse transcription. After hybridization, high stringency washing conditions were employed to remove nonspecific hybrids from the array. The remaining signal, emanating from each gene target, was measured using a Phosphorimager. Gene expression was reported as Phosphor Stimulating Luminescence (PSL) which reflects the level of phosphor signal generated from the probe hybridized to each of the gene targets represented on the array. A local background signal subtraction was performed before the total signal generated from each array was used to normalize gene expression between the different hybridizations. The value presented after "[array code]:" represents the mean of the duplicate values, following background subtraction and probe normalization. One of skill in the art could routinely use this information to identify normal and/or diseased tissue(s) which show a predominant expression pattern of the corresponding polynucleotide of the invention or to identify polynucleotides which show predominant and/or specific tissue and/or cell expression.

[40] Column 9 in Table 1A provides a chromosomal map location for certain polynucleotides of the invention. Chromosomal location was determined by finding exact matches to EST and cDNA sequences contained in the NCBI (National Center for Biotechnology Information) UniGene database. Each sequence in the UniGene database is assigned to a "cluster"; all of the ESTs, cDNAs, and STSs in a cluster are believed to be derived from a single gene. Chromosomal mapping data is often available for one or more

sequence(s) in a UniGene cluster; this data (if consistent) is then applied to the cluster as a whole. Thus, it is possible to infer the chromosomal location of a new polynucleotide sequence by determining its identity with a mapped UniGene cluster.

- [41] A modified version of the computer program BLASTN (Altshul et al., J. Mol. Biol. 215:403-410 (1990); and Gish and States, Nat. Genet. 3:266-272 (1993)) was used to search the UniGene database for EST or cDNA sequences that contain exact or near-exact matches to a polynucleotide sequence of the invention (the 'Query'). A sequence from the UniGene database (the 'Subject') was said to be an exact match if it contained a segment of 50 nucleotides in length such that 48 of those nucleotides were in the same order as found in the Query sequence. If all of the matches that met this criteria were in the same UniGene cluster, and mapping data was available for this cluster, it is indicated in Table 1A under the heading "Cytologic Band". Where a cluster had been further localized to a distinct cytologic band, that band is disclosed; where no banding information was available, but the gene had been localized to a single chromosome, the chromosome is disclosed.
- Once a presumptive chromosomal location was determined for a polynucleotide of the invention, an associated disease locus was identified by comparison with a database of diseases which have been experimentally associated with genetic loci. The database used was the Morbid Map, derived from OMIM<sup>TM</sup> (supra). If the putative chromosomal location of a polynucleotide of the invention (Query sequence) was associated with a disease in the Morbid Map database, an OMIM reference identification number was noted in column 10, Table 1A, labelled "OMIM Disease Reference(s)". Table 5 is a key to the OMIM reference identification numbers (column 1), and provides a description of the associated disease in Column 2.

## TABLE 1B

Clone ID NO:Z	SEQ ID NO:X	CONTIG ID:	BAC ID: A	SEQ ID NO:B	EXON From-To
HE2KJ64	12	906019	AC020570	659	1-67
				İ	406-542
					1507-1624
					2333-2429
		·			4080-4222
					4398-4455
					4561-4630
					4836-4971
					7386-7427
	j			1	7521-7596
HE2KJ64	12	906019	AC020570	660	1-247
HLICC37	14	856958	AL365356	661	1-195
					1135-2232
		Ì			2239-3110
HLICC37	14	856958	AL365356	662	1-173
HLICC37	14	856958	AL365356	663	1-141
HLTER04	23	590990	AC018845	664	1-273
					320-800
		ĺ			866-1324
					1551-2419
					3945-4348
					5055-5373
•					5597-5685
	•	1			6123-6519
					7020-7482
		1			7751-7856
			·		8955-9162
					9398-9496
					10809-11159
					13498-13544
					13809-14276
					14343-14490
					14632-14762
					16544-18402
HLTER04	23	590990	AC007338	665	1-273
				Ì	320-801
					867-1325
	1			1	1552-2420

					3946-4349
			}		5056-5374
					5598-5686
					6124-6520
					7021-7483
			1		7752-7857
ı					8956-9163
					9399-9497
		1			10810-11160
					13499-13545
1			İ		13810-14277
					14344-14491
		ļ			14633-14763
					16545-18403
HLTER04	23	590990	AC018845	666	1-249
HLTER04	23	590990	AC007338	667	1-249
H2MBY83	25	752124	AC017104	668	1-540
H2MBY83	25	752124	AC017104	669	1-548
HMZAD58	27	975304	AC078916	670	1-364
HMZAD58	27	975304	AC022305	671	1-686
HMZAD58	27	975304	AC002518	672	1-247
HMZAD58	27	975304	AC072032	673	1-364
HMZAD58	27	975304	AC078916	674	1-288
HMZAD58	27	975304	AC072032	675	1-288
HCHNH17	28	975378	AC026236	676	1-141
HBIMF04	36	951601	AL022328	677	1-103
					1215-1770
	·				2471-2545
					3028-3108
					3680-3960
					4352-4494
				1	4925-5476
					6623-6828
					6888-9053
					9409-10241
HBIMF04	36	951601	AL022328	678	1-333
HBIMF04	36	951601	AL022328	679	1-186
					376-570
	1				1511-2312
					2355-2630
			<u> </u>		2996-3446

		<u> </u>	·		3617-4004
		j			4225-5042
	i				5275-5664
£	İ	1			5695-5783
]	}		J		6915-7130
	•			·	7265-7787
			•		8377-9065
					9159-9294
				!	9608-9952
					10071-10419
					11431-11799
				i	12322-12621
1		1			12641-12911
					14491-14580
					14653-14848
					15670-15856
					15949-16109
		1			16183-16596
HOCQD08	39	972981	AC018568	680	1-1718
HOCQD08	39	972981	AC018568	681	1-425
HE8DL23	43	693641	AL135999	682	1-63
			·	·	405-942
		ł	1		1196-1502
			·		2152-6417
		į			6659-6755
					7033-7385
	Ì				7481-7535
	İ				7647-8163
					8230-8492
					8590-9909
					10114-10360
	ļ		}		10420-10783
,		Ì			10970-11960
			1		12018-13492
					14130-14528
					14563-15789
HE8DL23	43	693641	AL135999	683	1-410
HAJBU67	55	856922	AC008910	684	1-1685
					1960-2928
HAJBU67	55	856922	AC026230	685	1-1686
					1961-2933

TTA IDIICA	155	056000	TA C000010	1000	1 200
HAJBU67	55	856922	AC008910	686	1-326
HAJBU67	55	856922	AC026230	687	1-91
HAJBU67	55	856922	AC026230	688	1-325
HCEMY90	68	932927	AC024242	689	1-274
					1243-1357
				<u></u>	1994-2270
HCEMY90	68	932927	AF214633	690	1-274
				1	1243-1357
					1994-2270
HCEMY90	68	932927	AC024242	691	1-232
HCEMY90	68	932927	AF214633	692	1-130
HHFLF63	69	933854	AC023295	693	1-75
					1512-1564
HDTDG41	72	942490	AL137848	694	1-175
					2422-2550
					3441-3583
					4018-4129
	:				8219-8689
					9767-9876
					11592-11892
					14228-14324
1					15025-15162
					16319-16590
					17309-18595
HDTDG41	72	942490	AL137848	695	1-196
HFEBN52	82	810429	AL136001	696	1-61
					290-371
					654-779
			·		2128-2223
					2337-2372
					2507-2674
		;			3747-4249
					4554-4644
					5223-5557
,					5604-5916
					6827-6930
					6949-7329
					7852-8047
HFEBN52	82	810429	AL359399	697	1-61
					290-371
					654-779

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					2128-2223
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					2507-2674
					3747-4249
					4554-4644
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					6825-6928
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	·				6757-7115
					8075-8329
		·			8778-8876
					12305-12451
	* .				13119-13275
					16208-17104
HNSMB24	235	971537	AC015555	768	1-674
HNSMB24	235	971537	AP001623	769	1-674

Table 1B summarizes additional polynucleotides encompassed by the invention [43] (including cDNA clones related to the sequences (Clone ID NO:Z), contig sequences (contig identifier (Contig ID:) contig nucleotide sequence identifiers (SEQ ID NO:X)), and genomic sequences (SEQ ID NO:B). The first column provides a unique clone identifier, "Clone ID NO:Z", for a cDNA clone related to each contig sequence. The second column provides the sequence identifier, "SEQ ID NO:X", for each contig sequence. The third column provides a unique contig identifier, "Contig ID:" for each contig sequence. The fourth column, provides a BAC identifier "BAC ID NO:A" for the BAC clone referenced in the corresponding row of the table. The fifth column provides the nucleotide sequence identifier, "SEQ ID NO:B" for a fragment of the BAC clone identified in column four of the corresponding row of the table. The sixth column, "Exon From-To", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence of SEQ ID NO:B which delineate certain polynucleotides of the invention that are also exemplary members of polynucleotide sequences that encode polypeptides of the invention (e.g., polypeptides containing amino acid sequences encoded by the polynucleotide sequences delineated in column six, and fragments and variants thereof).

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W O 01/	33320													, 0501,	01047
NT To		445		232	277	364		243	121	151	449			125	267
NT From		212		86	35	2		19	41	53	3			3	175
Score/ Percent	Identity	82%		19.62	6.99	74%		100%	15.79	33.1	%6L			65.2	6'96
PFam/NR Accession Number		gi 4062512 dbj BAA3 5699.1		PF00419	PF00188	gi 4324682 gb AAD1	6986.1	sp AAF43006 AAF43 006	PF00023	PF00023	pir H65113 H65113			PF00480	PF00023
PFam/NR Description		Hypothetical fimbrial chaperone in pepN-pyrD	intergenic region. [Escherichia coli]	PFAM: Fimbrial proteins	PFAM: SCP-like extracellular Proteins	(AF109674) late gestation	lung protein 1 [Rattus norvegicus]	Diacylglycerol kinase iota (Fragment).	PFAM: Ank repeat	PFAM: Ank repeat	hypothetical 30.8 kD	intergenic region -	Escherichia coli (strain K-12)	PFAM: ROK family	PFAM: Ank repeat
Analysis	Method	blastx.14		HMMER 1.8	HMMER	blastx.14		blastx.2	HMMER 1.8	HMMER 2.1.1	blastx.2			HMMER 2.1.1	HMMER 2.1.1
SEQ ID	NO:X	11		237	12			13	238	14	15			239	16
Contig ID:		1106393		739539	906019			1178626	699372	826928	1121900			848220	823850
Clone ID	NO:Z	HFRBN59		HFRBN59	HE2KJ64	_		HAGDV32	HAGDV32	HLICC37	HBGBU96			HBGBU96	HAJCQ63

														<b>,</b> -		,								
628	322	382	384	448	419	256	146	450	144	533	66	66	44	468	295	556	561	308	323	203	353	323	323	323
242	245	230	33/	404	378	173	66	151	П	447	1	19	3	7	197	131	499	216	186	57	195	174	219	192
63%	17.86	62%	81%	%98	71%	36%	43%	91%	100%	%68	30%	46%	13.97	72%	62.3	72%	47%	59.5	41%	34%	37%	36%	48%	43%
sp Q9UQR3 Q9UQR 3	PF00023	gi 488505 dbj BAA06	418.1					gi 5712756 gb AAD4	7636.1 AF160798_1				PF00023	sp CAB89816 CAB8 9816	PF00023	gi 5262748 emb CAB	45688.1	PF00023	gi 790608 gb AAA85	854.1				
CENTAURIN BETA2.	PFAM: Ank repeat	similar to HUMORFU	(DZ6069) [Homo sapiens]					(AF160798) calcium	transporter CaT1 [Rattus	norvegicus]			PFAM: Ank repeat	Shank3b protein.	PFAM: Ank repeat	(AJ133120) Proline rich	synapse associated protein 2 [Rattus norvegicus]	PFAM: Ank repeat	UNC-44 [Caenorhabditis	elegans]				
blastx.2	HMMER 1.8	blastx.14						blastx.14					HMMER 1.8	blastx.2	HMMER 2.1.1	blastx.14		HMMER 2.1.1	blastx.14					
17	240						ļ	18					241	19	242			20						
1153903	926188							1096389					959139	1152327	903653			924647						
HLMMV66	HLMMV66							HLWAR08					HLWAR08	HBGTT76	HBGTT76		,	HMCF024						

323	182	173	167	182	323	200	170	71	71	65	462	574	757	482	962	467	388	1915	1804	1585	478	762	1041	350	244
174	102	06	09	75	219	78	78	3	3	6	427	476	479	291	786	285	62	2088	2097	1728	531	1037	1085	06	191
34%	48%	46%	38%	33%	45%	79%	29%	47%	34%	47%	20%	37.4	33%	31%	25%	29%	28%	53.39	%99	45%	%99	. 63%	<del> </del> %09	128.3	12.82
						•						PF00023	gb AAC96986.1				sp AAF67491 AAF67 491	PF00412	gi 841318 gb AAA85	718.1		gi 841318 gb AAA85	718.1	PF00684	PF00023
												PFAM: Ank repeat	contains 10 ankyrin-like	repeats; similar to human	ankyrin, 1 bursaria	Chlorella virus 1]	Sterol regulatory element binding protein 3.	PFAM: LIM domain containing proteins	mutant sterol regulatory	element binding protein-2	1	mutant sterol regulatory	element binding protein-2	PFAM: DnaJ central domain (4 repeats)	PFAM: Ank repeat
		,	-									HIMMER 2.1.1	blastx.2				blastx.2	HMMER 1.8	blastx.14			blastx.14		HMMER 2.1.1	HMMER 1.8
												21					22	243				244		23	245
												973137					1012465	892956				975276		290990	823859
												HBIOM94			00		HBJLR11	HBJLR11				HBJLR11		HLTER04	HMSMU30

493	493	1299	438	739	1813	2509	1972	1894	2104	1360	1546	1009		1009	1021	918	1239	1182	1182	1206	1185	1182	1188	609
80	80	130	340	362	368	1964	1712	1730	2000	1280	1445	968		116	917	820	91	163	130	163	166	106	103	163
137	28%	51%	35.4	196.5	%68	100%	%26	25%	34%	33%	73%	31.3		%06	34%	321.4	%56	39%	37%	34%	36%	32%	33%	39%
PF01951	sp Q9VD92 Q9VD92	sp BAA91856 BAA9 1856	PF00023	PF01412	gi 4691728 gb AAD2	8047.1 AF124491_1						PF00412		sp AAF34411 AAF34	411	PF00023	gb AAA64834.1							
PFAM: Protein of unknown function	CG6353 PROTEIN.	CDNA FLJ10852 FIS, CLONE NT2RP4001498, WEAKLY SIMILAR TO 1	PFAM: Ank repeat	PFAM: Putative GTP-ase activating protein for Arf	(AF124491) ARF	GTPase-activating protein	GIT2 [Homo sapiens]				,	PFAM: LIM domain	containing proteins	LIM and cysteine-rich	domains protein 1.	PFAM: Ank repeat	ankyrin G [Homo sapiens]							
HMMER 2.1.1	blastx.2	blastx.2	HMMER 2.1.1	HIMMER 2.1.1	blastx.14			,				HMMER	2.1.1	blastx.2		HIMMER 2.1.1	blastx.2							
25		26	246	27								28				247								
752124		1164739	810424	975304								975378				971772								
H2MBY83		<b>НВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ</b> <b>ТВ ТВ ТВ ТВ ТВ ТВ ТВ ТВ</b>	HBUAH93	HMZAD58								HCHNH17				HBWAJ55								

597	244	440	1031	75	1074	519	1314	195	369	396	312	288	312	291	312	279	356	359	695	716	308	422	797	828	
148	1230	354	81	34	985	148	25	106		13	28	4	4	4	4	4	315	315	255	426	144	300	726	22	
38%	100%	85	100%	100%	79%	147.9	%66	72.3	%02	48%	44%	44%	39%	36%	35%	34%	57%	53%	26.4	23%	32%	46%	28%	132	
	sp Q9Y576 Q9Y576	PF00023	gi 5306062 gb AAD4	1894.1 AF156777 1	}	PF01980	sp BAA91013 BAA9 1013	PF00023	sp Q9VCA7 Q9VCA	7									PF01942	gi 3128208 gb AAC2	6688.1			PF02005	
	ASB-1 PROTEIN.	PFAM: Ank repeat	(AF156777) ASB-1	protein [Homo sapiens]		PFAM: Uncharacterised protein family	CDNA FLJ20206 FIS, CLONE COLF1582.	PFAM: Ank repeat	CG6268 PROTEIN.										PFAM: Domain of unknown function	(AC004077) hypothetical	protein [Arabidopsis	thaliana]		PFAM: N2,N2-	dimethylguanosine tRNA methyltransferase
	blastx.2	HMMER 2.1.1	blastx.14			HMMER 2.1.1	blastx.2	HMMER 2.1.1	blastx.2										HMMER 2.1.1	blastx.14				HMMER	2.1.1
	30	248				31	5 * *	32										٠	33					34	
	1152346	911597				919538		911586											926488					945055	
	HNJCE31	HNJCE31				HKAIU14		HCE4112		-		20							HFOY118					HHEDM89	٠

720	2310	2337	1043	1130	1249	237	,	207		327 339	391	1399	307	715	326
16	2104	529	336	405	1085	160		46		13 292	293	2	38	320	78
35%	65.2	%66	191.2	%66	%08	100%		67.2		67% 37%	50.4	%29	42.2	78%	%86
dbj BAA30948.1	PF00563	dbj BAA35528.1	PF01963	emb CAB63043.1				PF00412		gi 1710382 gb AAB3 8287.1	PF00023	sp 097902 097902	PF02000	pir A61382 A61382	
(AP000007) 381aa long hypothetical N2,N2- dimethylguanosine tRNA methyltransferase [Pyrococcus horikoshii]	PFAM: Domain of unknown function 2	Hypothetical 67.7 kd protein CY02B10.18C. [Escherichia coli]	PFAM: TraB family	(AL022328) dJ402G11.4	(novel protein similar to	C. elegans F38A5.2	(isoform 1)) [Homo	PFAM: LIM domain	containing proteins	ajuba; jub [Mus musculus]	PFAM: Ank repeat	DIFFERENTIATION ENHANCING FACTOR 1.	PFAM: Protein of unknown function	phosphorylation	regulatory protein HP-10 - human
blastx.2	HMMER 2.1.1	blastx.2	HMMER 2.1.1	blastx.2				HMMER	2.1.1	blastx.14	HMMER 2.1.1	blastx.2	HMMER 2.1.1	blastx.14	·
	35		36					249		·	38	<del> </del>	39		
	945288		951601		_			946972			823854		972981		
	HFXKW18		HBIMF04				191	HEEAU28			HDPKI66		НОСОДО8		

1517	401	192	322	319	343	406	351	357	51	456	585	223	340	159	460
75	330	19	2	2	29	29 408	178	22	22	581	662	155	146	109	164
%96	20.99	87.1	194.1	%56	131.4	83% 100%	43.25	36%	%06	64%	30%	28.1	%99	94%	150.3
sp BAA91856 BAA9 1856	PF00023	PF00635	PF01602	pir A36680 A36680	PF01602	sp 075504 075504	PF00412	sp O70209 O70209		gi 1905874 gb AAC0	5580.1	PF00035	sp 075569 075569		PF01217
CDNA FLJ10852 FIS, CLONE NT2RP4001498, WEAKLY SIMILAR TO	PFAM: Ank repeat	PFAM: MSP (Major sperm protein) domain	PFAM: Adaptin N terminal region	gamma-adaptin precursor - mouse	PFAM: Adaptin N terminal region	GAMMA2-ADAPTIN.	PFAM: LIM domain containing proteins	ALPHA-ACTININ-2	ASSOCIATED LIM PROTEIN.	carboxyl terminal LIM	domain protein [Homo sapiens]	PFAM: Double-stranded RNA binding motif	PROTEIN ACTIVATOR	OF THE INTERFERON- INDUCED PROTEIN KINASE	PFAM: Clathrin adaptor
blastx.2	HMMER 1.8	HMMER 2.1.1	HIMMER 2.1.1	blastx.2	HMMER 2.1.1	blastx.2	HMMER 1.8	blastx.2		blastx.14		HIMMER 2.1.1	blastx.2		HMMER
40	250.	41	42	·	43		44			251		45			46
1228283	502892	609468	909029		693641		928851			948605		775616			715097
HDPRP54	HDPRP54	HE2BW32	HAJAU21		HE8DL23		HFTCM92			HFTCM92		HE6BQ76			HAMFP60

	305	403	390	217	399		249	833	833	815	725	638	737	707	909	710	593	731	503	413	521	707	404	410	969	827
	129	846	283	2	211		190	735	126	126	123	123	339	111	219	129	201	204	144	120	123	342	135	117	501	744
	78.2	36%	26.1	%88	 %99		17.29	187.9	33%	32%	30%	31%	36%	30%	37%	73%	36%	25%	38%	36%	31%	32%	41%	73%	32%	42%
	PF01217	7V6V6V7 Q9V9V7	PF00035	sp BAA91862 BAA9	1862		PF00023	PF00023	gi 557084 gb AAC37	208.1																
complex small chain	PFAM: Clathrin adaptor complex small chain	CG1800 PROTEIN.	PFAM: Double-stranded RNA binding motif	CDNA FLJ10860 FIS,	CLONE NT2RP4001568,	WEAKLI SHMILAK IO 1	PFAM: Ank repeat	PFAM: Ank repeat	ankyrin [Drosophila	melanogaster]																
2.1.1	HMMER 2.1.1	blastx.2	HMMER 2.1.1	blastx.2			HMMER 1.8	HMMER 2.1.1	blastx.14													,				
	47	48	252	49		·	253	254																		
	715098	1150900	859840	1165338			944518	904598																		
	HHFHY84	HE6FD03	HE6FD03	HDTFT90			HDTFT90	HPJCU63			•															

																		_									
815	833	737	836	824	970	812	349		352	136				246	1135				712	595	355	508	557		909	1458	278
726	738	009	732	732	923	726	9/1		68	14				148	2				443	2	257	2	525		514	562	3
40%	37%	32%	78%	35%	%95	34%	47.4		95%	85%				46.4	%66				102.6	61%	93.8	%66	%06		28.1	82%	%96
							PF01454		sp O76058 O76058					PF01602	sp BAA91511 BAA9	1511			PF00636	sp Q9Z1P7 Q9Z1P7	PF00023	gi 1136404 dbi BAA1	1489.1		PF01605	pir T34532 T34532	
							PFAM: MAGE family		DJ1409.2	(MELANOMA-	ASSOCIATED	ANTIGEN MAGE	LIKE).	PFAM: Adaptin N terminal region	CDNA FLJ10259 FIS,	CLONE	HEMBB1000947,	HIGHLY SIMILAR 10 I	PFAM: RNase3 domain.	NG28.	PFAM: Ank repeat	similar to ankyrin of	Chromatium vinosum.	[Homo sapiens]	PFAM: eRF1-like proteins	hypothetical protein	DKFZp434B1517.1 -
							HMMER	2.1.1	blastx.2					HMMER	blastx.2				HIMMER 2.1.1	blastx.2	HIMMER	blastx.14			HMMER 2.1.1	blastx.2	
							51							22	53				255	54	256				55	56	
							793203							796509	1158800				914398	1197903	932013				856922	1204696	
							HFITE38							нDPDH64	R HFKKS58				HFKKS58	HE8CM38	HE8CM38				HAJBU67	HHEHD10	

***	Λ.	/55326	

PFAM: LIM domain containing proteins PFAM: Double-stranded RNA binding motif ASB-3 PROTEIN (CDNA FLJ10123 FIS, CLONE HEMBA1002939, WEAKLY 1 PFAM: Ank repeat (AF156778) ASB-3 protein [Homo sapiens] PFAM: Ank repeat WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). hypothetical protein DKFZp434B1517.1 - human (fragment) PFAM: LIM domain containing proteins Hypothetical 35.8 kDa protein. PFAM: Putative GTP-ase activating protein for Arf	human (fragment)				
1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00   1.00	├	PF00412	28.72	48	224
1031960 58   blastx.2   ASB-3 PROTEIN (CDNA		PF00035	12.86	25	114
FLJ10123 FIS, CLONE   HEMBA1002939,   WEAKLY 1   2.1.1   blastx.14   (AF156778) ASB-3   protein [Homo sapiens]   protein [Homo sapiens]   protein [Homo sapiens]   blastx.2   WUGSC:H_DJ1035002.1   PROTEIN   PRAGMENT).   1091714   60   blastx.2   hypothetical protein   DKFZp434B1517.1 - human (fragment)   154785   61   blastx.2   Hypothetical 35.8 kDa   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein.   protein	<del>                                     </del>	sp Q9Y575 Q9Y575	82%	199	702
HEMBA1002939, WEAKLY 1 2.1.1 blastx.14 (AF156778) ASB-3 protein [Homo sapiens] HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 pROTEIN (FRAGMENT). HO91714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf 2.1.1 activating protein for Arf	FLJ10123 FIS, CLONE		94%	695	751
911594 258 HMMER PFAM: Ank repeat 2.1.1 blastx.14 (AF156778) ASB-3 protein [Homo sapiens]  911588 59 HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). 1091714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment) 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	HEMBA1002939, WEAKLY 1		42%	674	751
911588 59 HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). 1091714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	ER	PF00023	128.9	430	528
protein [Homo sapiens]  HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT).  1091714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	14	gi 5306064 gb AAD4	94%	199	786
911588 59 HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). (FRAGMENT). DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	protein [Homo sapiens]	1895.1 AF156778_1	37%	433	615
911588 59 HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). 1091714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf			10%	792	884
911588 59 HMMER PFAM: Ank repeat 2.1.1 blastx.2 WUGSC:H_DJ1035002.1 PROTEIN (FRAGMENT). (FRAGMENT). DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf			36%	718	783
blastx.2 WUGSC:H_DJ1035002.1		PF00023	47	160	258
PROTEIN (FRAGMENT).   PROTEIN (FRAGMENT).   1091714   60   blastx.2   hypothetical protein	-	gp[Q9UDM3]Q9UD	28%	169	402
1091714 60 blastx.2 hypothetical protein DKFZp434B1517.1 - human (fragment)  894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	PROTEIN (FRAGMENT).	M3	53%	438	554
894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf		pir T34532 T34532	95%	235	402
894375 259 HMMER PFAM: LIM domain 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	DKFZp434B1517.1 -		93%	685	77.1
894375 259 HMMER PFAM: LIM domain 1.8 containing proteins 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	human (fragment)		51%	27	128
894375 259 HMMER PFAM: LIM domain 1.8 containing proteins 1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf			25%	611	169
1154785 61 blastx.2 Hypothetical 35.8 kDa protein. 2 909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf		PF00412	26.8	133	252
909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf	1	sp CAC09448 CAC0	%66	8	811
909634 260 HMMER PFAM: Putative GTP-ase 2.1.1 activating protein for Arf		9448			
activating protein for Arf		PF01412	184.7	256	819
	$\dashv$				
blastx.14   (AL031633) similar to   gi 3880859 emb CA	$\dashv$	gi 3880859 emb CAA	21%	265	510

732	861 172	379	664	486	483	489	544	396	477	726	504	486	504	405	456	192	366	357	372	372	357	366	381	84	360
502	733	. 2	2	199	208	181	488	298	52.	52	4	7	7	7	4	94	106	97	103	100	103	103	196	1	37
42%	46%	104.3	100%	28%	36%	35%	21%	47.6	40%	33%	35%	32%	31%	34%	27%	63.1	42%	36%	35%	35%	38%	31%	40%	35%	40%
21032.1		PF00637	gi 969024 gb AAC48 524.1	sp BAA91302 BAA9	1302			PF00023	pir T18184 T18184		-					PF00023	gi[2447128 gb AAC9	6986.1		-			•		sp 000151 CL36_HU MAN
Ank repeat (2 domains);	cDNA 1	PFAM: 7-fold repeat in Clathrin and VPS	clathrin heavy chain [Bos taurus]	CDNA FLJ20636 FIS,	CLONE KAT03434.			PFAM: Ank repeat	ankyrin repeat protein	A682L - Chlorella virus	PBCV-1					PFAM: Ank repeat	contains 10 ankyrin-like	repeats; similar to human	1 Paramecium bursaria	Chlorella virus 1]	ı				LIM DOMAIN PROTEIN CLP-36.
		HMMER 2.1.1	blastx.14	blastx.2		<del>.</del>		HMMER 2.1.1	blastx.2							HMMER 2.1.1	blastx.14								blastx.2
		62		63				261	64							262									99
		914810		1117857				810432	1154798							911592									1181020
		HOHAS44		HE80F42				HE80F42	HSKHS71							HSKHS71									HISBT75

302	308	179	2780	1748	1583	3705	1940	3720	1892	3461	3248	3128	3514	1937	408	456	456	234	585	605	654		285	654	336
129	183	12	3205	3718	1741	3788	2014	3788	1936	3535	3289	3196	3558	2011	340	115	115	<i>L</i> 9	19	546	487		4	421	777
43.09	64%	25%	642.4	%86	100%	100%	36%	36%	46%	48%	21%	30%	46%	28%	30.1	48%	39%	48.7	62%	65%	110.2		36%	44%	35%
PF00412	gi 1905874 gb AAC0	5580.1	PF00637	gi 969024 gb AAC48	524.1										PF00560	emb CAA73132.1		PF00855	sp O96028 O96028		PF02134		gi 54058 emb CAA44	465.1	
PFAM: LIM domain containing proteins	carboxyl terminal LIM	domain protein [Homo sapiens]	PFAM: 7-fold repeat in Clathrin and VPS	clathrin heavy chain [Bos	taurus]										PFAM: Leucine Rich Repeat	hypothetical protein	[Silene latifolia]	PFAM: PWWP domain	WHSC1 PROTEIN.		PFAM: Repeat in	ubiquitin-activating (UBA) proteins	Sbx [Mus musculus]	,	
HMMER 1.8	blastx.14		HMMER 2.1.1	blastx.14											HMMER 2.1.1	blastx.2		HMMER 2.1.1	blastx.2		HMMER	2.1.1	blastx.14		
263			99												<i>L</i> 9			89			69				
963281			930964												931402			932927	~		933854				
HISBT75			HFVKF77										07		HJABW64			HCEMY90			HHFLF63				

70 HWWER PEAM: I envine Rich
2.1.1
blastx.14 similar to yeast adenylate cyclase (S56776) [Homo sapiens]
264 HMMER PFAM: Laminin B 1.8 (Domain IV)
72 HMMER PFAM: Leucine Rich 2.1.1 Repeat
blastx.14 proteoglycan I precursor
[Homo sapiens]
73 HMMER PFAM: Double-stranded 1.8 RNA binding motif
blastx.2 testis nuclear RNA
binding protein - mouse
74 blastx.2 ankyrin-related protein
unc-44 - Caenorhabditis
elegans (fragment)
265 HMMER PFAM: Ank repeat
+
blastx.2
75   blastx.2   probable ATPase SKD3
[imported] - mouse
2.1.1 PFAM: Ank repeat